# C py for the Elected Office (EO/P) PATENT COOPERATION TREATY

From the INTERNATIONAL BUREAU

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101	10:			
NOTIFICATION OF THE RECORDING OF A CHANGE  (PCT Rule 92bis.1 and Administrative Instructions, Section 422)  Date of mailing (day/month/year) 03 April 2002 (03.04.02)	KIM, Won-Ho 7th Fl., Teheran Bd. 825-33 Yoksam-dong, Kangpan ku Seoul 135-080 RÉPUBLIQUE DE CORÉE JUN 0 4 2007			
Applicant's or agent's file reference	IMPORTANT NOTIFICATION			
OPP000458KR	INFORTANT NOTIFICATION			
International application No. PCT/KR00/00664	International filing date (day/month/year) 26 June 2000 (26.06.00)			
The following indications appeared on record concerning:      X the applicant      X the inventor	the agent the common representative			
Name and Address	State of Nationality State of Residence			
SON, Seog-Jae JANG, Ḥa-Soo	KR KR Telephone No.			
	Facsimile No.			
	Teleprinter No.			
	i i			
The International Bureau hereby notifies the applicant that the X the person the name the add				
Name and Address	State of Nationality State of Residence			
SON, Seog-Jae JANG, Ha-Soo	KR KR Telephone No.			
	SECTION			
	Facsimile No.			
	Teleprinter No.			
3. Further observations, if necessary: Corrected version of IB/306(12.12.01). The perso applicant/inventor for US only.	n in box No. 2 will be the the designated Offices concerned			
4. A copy of this notification has been sent to:	(C)			
X the receiving Office	the designated Offices concerned			
the International Searching Authority	X the elected Offices concerned			
the International Preliminary Examining Authority	other:			
The International Durana (1980)	Authorized officer			
The International Bureau of WIPO 34, chemin des Colombettes	Ki-Nam HA			
1211 Geneva 20, Switzerland  Facsimile No : (41-22) 740 14 35	elephone No.: (41-22) 338.83.38			

	From the INTERNATIONAL BUREAU				
PCT	То:				
NOTIFICATION OF THE RECORDING OF A CHANGE  (PCT Rule 92bis.1 and Administrative Instructions, Section 422)  Date of mailing (day/month/year) 12 December 2001 (12.12.01)	KIM, Won-Ho 7th Fl., Teheran Bd. 825-33 Yoksam-dong, Kangnam-ku Seoul 135-080 RÉPUBLIQUE DE CORÉE				
Applicant's or agent's file reference OPP000458KR		IMPORTANT NOT	TIFICATION		
International application No.		nal filing date (day/month/y	/ear)		
PCT/KR00/00664	26 J	une 2000 (26.06.00)	·		
The following indications appeared on record concerning:      The applicant the inventor	the agei	nt the comm	on representative		
Name and Address		State of Nationality  KR	State of Residence KR		
SON, Seog-Jae JANG, Ha-Soo		Telephone No.	l vu		
		reteprione ivo.			
	•	Facsimile No.			
		Teleprinter No.			
2. The International Bureau hereby notifies the applicant that the	ne following	change has been recorded	concerning:		
the person the name the add	lress	the nationality	the residence		
Name and Address		State of Nationality	State of Residence		
·		KR	KR		
		Telephone No.			
		Facsimile No.	•		
		Teleprinter No.			
3. Further observations, if necessary:		• "	-		
4. A copy of this notification has been sent to:					
X the receiving Office	[	the designated Offices	s concerned		
the International Searching Authority	[	X the elected Offices cor	ncerned		
the International Preliminary Examining Authority		other:			
T	Authorized	officer			
The International Bureau of WIPO 34, chemin des Colombettes 1211 Geneva 20, Switzerland		Ning XU			
Facsimile No.: (41-22) 740.14.35	Telephone	No.: (41-22) 338.83.38			



To:

### From the INTERNATIONAL BUREAU

### **PCT**

### **NOTIFICATION OF ELECTION**

(PCT Rule 61.2)

Commissioner
US Department of Commerce
United States Patent and Trademark
Office, PCT
2011 South Clark Place Room
CP2/5C24
Arlington, VA 22202

in its capacity as elected Office		
Applicant's or agent's file reference		
OPP000458KR		
Priority date (day/month/year)		
26 June 1999 (26.06.99)		
	Applicant's or agent's file reference OPP000458KR  Priority date (day/month/year)	

1.	The designated Office is hereby notified of its election made:
	X in the demand filed with the International Preliminary Examining Authority on:
	16 January 2001 (16.01.01)
	in a notice effecting later election filed with the International Bureau on:
•	The sleeting [V]
2.	The election X was was not
	made before the expiration of 19 months from the priority date or, where Rule 32 applies, within the time limit under Rule 32.2(b).

The International Bureau of WIPO 34, chemin des Colombettes 1211 Geneva 20, Switzerland Authorized officer

Zakaria EL KHODARY

Telephone No.: (41-22) 338.83.38

Facsimile No.: (41-22) 740.14.35



### **PCT**

### **NOTIFICATION CONCERNING** AMENDMENTS OF THE CLAIMS

(PCT Rule 62 and Administrative Instructions, Section 417) From the INTERNATIONAL BUREAU

International filing date (day/month/year)

To:

Korean Intellectual Property Office 920 Dunsan-dong, So-gu 302-701 Daejeon Metropolitan City RÉPUBLIQUE DE CORÉE

in its capacity as International Preliminary Examining Authority

26 June 2000 (26.06.00)

Date of mailing (day/month/year) 22 February 2001 (22.02.01)

International application No.

PCT/KR00/00664

**Applicant** 

LG CHEMICAL LTD. et al

The International Bureau hereby informs the International Preliminary Examining Authority that no amendments under Article 19 have been received by the International Bureau (Administrative Instructions, Section 417).

> The International Bureau of WIPO 34, chemin des Colombettes 1211 Geneva 20, Switzerland

Authorized officer

Zakaria EL KHODARY

Facsimile No. (41-22) 740.14.35

Telephone No. (41-22) 338.83.38

### □ APPLICANT

### NOTIFICATION OF CHANGE OF

**⊠INVENTOR** 

To: Commissioner of

the Korean Industrial Property Office												
International Application No.			T/KR	00/00	664							
Name		LG Che	emical Li			Resid Reg.	No.		Countr Nation	ality	K	blic of orea
Applicant	Address	LG Twin Towe	er, Yo	ido-don	g 20,				Seou.	1, 1	50-721, F	Republic of
e termina e .	Name	KIM, Won	-Но	Agent	t's Code		199 <u>8</u> 0023		Tel. No	s:	+82-2-3	3458-070
Agent	Address	Teheran Bd., Fax: +82-2-5			m-dong	, Kanç email@	nam- youm	ku, S	Seoul,	Repu	ablic of F	Corea
Cause of Cl	nange	Addition of	an in	ventor	(SON,	Seog	-Jae	e, an	d JAN			
		Name		G, Hea-			Ì	Reside Reg. N	lo.	N:	ationality	Republic of Korea
	Former	Address	Deokil Hanmaeum Apt. 101-1205, Wooam-dong, Sangdang-ku, Cheongju city, Choongcheongbuk-do, 360-200, Republic of Korea									
		Name	SON	, Seog	-Jae	·		idence . No.		Cour Natio	ntry of onality	Republic of Korea
,		Address	LG cit	Sataek y 680-	A-404 B32, R	, Shi epubl	njec ic c	ong 2 of Ko	-dong rea	, 11	82, Nam-	ku, Ulsan-
Contents of	-	Designated	Stat	e(s)	US or	ıly				,		<del>, </del>
Notifi- cation		Name	1	JANG, Ha-Soo			Reg	idence , No.		N	lationality	Republic of Korea
	New	Address	Shi Seo	Shindonga Apt. 3-403, Yeomchang-dong 289, Kangseo-Seoul, 157-864, Republic of Korea					(angseo-ku,			
	1	Designated	Stat	e(s)	US or	nly						
		Name	KAN	IG, Hea			Reg	idence g. No.	- }	N	Country of Vationality	of Korea
		Address	Dec Che Kor	ongju	nmaeum city,	Apt. Choor	10: ngche	1-120 eongb	)5, Wo	oam- , 36	-dong, Sa 50-200, F	ingdang-ku, Republic of
		Designated	Stat	e(s)	US or	ıly						

Submitted hereby is a notification pursuant to Article 82 of the Enforcement Regulations of the Patent Law.

Date(day/month/year)

07 November 2001 (07.11.2001)

(Seal)

KIM, Won-Ho Applicant (Agent)

### Attached Document(s):

- A copy of the document(s) substantiating the contents of notification (in case of notification 1. by new applicant)
- A certified copy of seal impression (in case of notification by new applicant)
- A copy of the document(s) substantiating the power of attorney, if any

### **⊠APPLICANT**

### NOTIFICATION OF CHANGE OF

Address

Address

Address

Name

Name

New

Designated

Designated

State(s)

State(s)

Korea

Designated State(s)

JANG, Ha-Soo

Shindonga Apt.

KANG, Hea-Chun

#### □INVENTOR

Reg. No.

Residence

Reg. No.

Residence

Deokil Hanmaeum Apt. 101-1205, Wooam-dong, Sangdang-ku, Cheongju city, Choongcheongbuk-do, 360-200, Republic of

Reg. No.

city 680-832, Republic of Korea US only

Seoul, 157-864, Republic of Korea

US only

US only

LG Sataek A-404, Shinjeong 2-dong, 1182, Nam-ku, Ulsan-

Republic

of Korea

of Korea

Country of | Republic

Country of

Nationality

Nationality

3-403, Yeomchang-dong 289, Kangseo-ku,

To: Commissioner of the Korean Industrial Property Office PCT/KR00/00664 International Application No. Republic of Residence Country of Name LG Chemical Ltd. et al. Korea Reg. No. Nationality Applicant Republic of LG Twin Tower, Yoido-dong 20, Yongdungpo-ku, Seoul, 150-721, Address Korea 9-1998-+82-2-3458-070 Agent's Code Tel. No. KIM, Won-Ho Name 000023-8 Agent Teheran Bd., 825-33, Yoksam-dong, Kangnam-ku, Seoul, Republic of Korea Address Fax: +82-2-553-5254 E-mail: email@youme.com Cause of Change Assignment Republic Residence Country of LG Chemical Ltd. Name of Korea Nationality Reg. No. LG Twin Tower, Yoido-dong 20, Yongdungpo-ku, Seoul, 150-721, Address Republic of Korea Country of Republic Residence SON, Seog-Jae Name Nationality of Korea Reg. No. 1182, Nam-ku, Ulsan-LG Sataek A-404, Shinjeong 2-dong, Address city 680-832, Republic of Korea of Republic Residence Country Former JANG, Ha-Soo Name Nationality of Korea Reg. No. Yeomchang-dong 289, Kangseo-ku, 3-403, Shindonga Apt. Address Republic of Korea Seoul, 157-864, Republic Residence Country of KANG, Hea-Chun Name of Korea Nationality Reg. No. Deokil Hanmaeum Apt. 101-1205, Wooam-dong, Sangdang-ku, Cheongju city, Choongcheongbuk-do, 360-200, Republic of Address Korea Republic Residence Country of LG Chemical Co., Ltd. Name Nationality of Korea Reg. No. Contents of LG Twin Tower, Yoido-dong 20, Youngdungpo-ku, Seoul, 150-721, Notifi-Address Republic of Korea cation All Designated States except US State(s) Designated Republic Country of | Residence SON, Seog-Jae Name Nationality of Korea

### Submitted hereby is a notification pursuant to Article 82 of the Enforcement Regulations of the Patent Law.

Date(day/month/year)

13 November 2001 (13.11.2001)



Applicant (Agent)

KIM, Won-Ho

(Seal)

### 

- 1. A copy of the document(s) substantiating the contents of notification (in case of notification by new applicant)
- 2. A certified copy of seal impression (in case of notification by new applicant)
- 3. A copy of the document(s) substantiating the power of attorney, if any

6.7" " 1 1 " × 78

### **APPLICANT**

### NOTIFICATION OF CHANGE OF

□INVENTOR

To: Commissioner of the Korean Industrial Property Office

al Application	n No. PC'	T/KR00/00664		<del></del>		e Popubi	ic of	
Name	\ LG Che		Reg. No.	N	ationalit	v Ko	cea	
Address		r, Yoido-dong 20,		· · · · ·	Seour,		·	
Name	KIM, Won-		00002	3-8 1				
Address	Fax: +82-2-55	Fax: +82-2-553-5254 E-Mail: email:						
hange	Change of t	he designated Sta	tes of tw	o of t	he app	licants	Republic	
	Name	I.G. Chemical I.td.		Residen	ce	Nationality 01	of Korea	
	Address .	LG Twin Tower, 'Republic of Kore	Yoido-dong a				Republic	
	Name	SON, Seog-Jae		Reg. No		Nationality	of Korea	
	Address	LG Sataek A-40 city 680-832,	4, Shinje Republic	of Kor	ea			
Former	Name	JANG, Ha-Soo		Dog M	.	Nationality	of Korea	
	Address	Seoul, 157-864, Republic of Roled						
	Name	KANG Hea-Chun	KANG, Hea-Chun			Nationality	of Korea	
	Address	Cheongju city,	m Apt. 10 Choongcl	01-1205 neongbu	nk-do,		<u>.</u>	
	Name	LG Chemical Ltd.	Reg. N	lo.		Nationality	Republic of Korea	
	Address	LG Twin Tower, Yoido-dong 20, Yongdungpo-ku, Seoul, 150-72. Republic of Korea						
	Designate	d State(s) All			tes ex	cept US	Republi	
	Name	SON, Seog-Jae	g	ea No		Nationality	of Kore	
·	Address	city 680-832,	Republic	eong 2 of Ko	-dong, rea	1182, Nam-	ku, Ulsan	
	Designate	d State(s) US		) i d o = 00		Country of	Republi	
New	Name	JANG, Ha-Soo	l r	Rea No.	1	Nationality	of Kore	
	Address	Seoul, 157-86	Seoul, 157-864, Republic of Korea					
1	Designat	ed State(s)   US		Didana		Country of	Republi	
	Name	KANG, Hea-Chu	ın j	Dea No	ì	Nationality	of Kore	
	Address	Deokil Hanmae Cheongju city Korea	eum Apt. y, Choong	101-120 cheong	05, Woodk-do	oam-dong, S	Republic (	
	Name Address Name Address Change	Name  LG Che Address  Name  KIM, Won- Address  Teheran Bd., Fax: +82-2-55  Change  Change of t  Name  Address  Designate  Name	Name  LG Chemical Ltd. et al.  Address Korea  Name  KIM, Won-Ho  Agent's Code  Address  Teheran Bd., 825-33, Yoksam-dong Fax: +82-2-553-5254  Change  Change of the designated State  Name  LG Chemical Ltd.  Address  Republic of Kore  Name  Address  LG Sataek A-40 city 680-832,  Name  Name  Name  LG Chemical Ltd.  Address  Former  Name  Address  Shindonga Apt. Seoul, 157-864  Name  Address  Name  LG Chemical Ltd.  LG Twin Tower, Republic of Kore  Address  Cheongju city, Korea  Name  LG Chemical Ltd.  LG Twin Tower, Republic of Kore  Address  Designated State(s)  Name  Address  LG Sataek A-4 city 680-832,  Designated State(s)  Name  Address  Designated State(s)  New  Name  JANG, Ha-Soo  Address  Shindonga Apt. Seoul, 157-864  Shindonga Apt. Seoul, 157-864  Shindonga Apt. Seoul, 157-864  Cheongju city, Korea  New  Name  Address  Designated State(s)  Name  KANG, Hea-Chu  Designated State(s)  New  Name  Address  Cheongju city, Korea  Cheongju city, Korea  New  Name  Address  Cheongju city, Korea  New  Name  Address  Cheongju city, Korea  New  Name  Address  Cheongju city, Korea  Deokil Hanmae, Cheongju city, Korea  Cheonglu cheongju city, Korea  Cheonglu ch	Name    LG Chemical Ltd. et al.   Residence Reg. No.	Name	Name LG Chemical Ltd. et al. Residence Reg. No. Nationality (Country of Nationality Korea)  Name KIM, Won-Ho Agent's Code 9-1998-000023-8  Name KIM, Won-Ho Agent's Code 9-1998-000023-8  Teheran Bd., 825-33, Yoksam-dong, Kangnam-ku, Seoul, Refax: +82-2-553-5254 E-mail: email@youme.com  Change of the designated States of two of the app Reg. No. LG Twin Tower, Yoido-dong 20, Yongdung Republic of Korea  Name LG Chemical Ltd. Residence Reg. No. LG Sataek A-404, Shinjeong 2-dong, City 680-832, Republic of Korea  Name JANG, Ha-Soo Residence Reg. No. Seoul, 157-864, Republic of Korea  Name KANG, Hea-Chun Residence Reg. No. Cheongju city, Choongcheongbuk-do, Korea  Name LG Chemical Ltd. Residence Reg. No. LG Twin Tower, Yoido-dong 20, Yongdung Republic of Korea  Name LG Chemical Ltd. Residence Reg. No. LG Twin Tower, Yoido-dong 20, Yongdung Republic of Korea  Name LG Chemical Ltd. Residence Reg. No. LG Twin Tower, Yoido-dong 20, Yongdung Republic of Korea  Designated State(s) All Designated States expended to Korea  Designated State(s) IDS only  New Name JANG, Ha-Soo Residence Reg. No. Residence Reg. No. Republic of Korea  Designated State(s) US only  Name JANG, Ha-Soo Residence Reg. No. Residence Re	Name	

## Submitted hereby is a notification pursuant to Article 82 of the Enforcement Regulations of the Patent Law.

Date(day/month/year)

07 November 2001 (07.11.2001)

Applicant (Agent)

KIM, Won-Ho

(Seal)



### 

1. A copy of the document(s) substantiating the contents of notification (in case of notification by new applicant)

2. A certified copy of seal impression (in case of notification by new applicant)

3. A copy of the document(s) substantiating the power of attorney, if any

### Power of Attorney

Agent (Common Representative)

Name: Patent Attorney KIM, Won-Ho

Address: Teheran Bldg., 825-33, Yoksam-dong,

Kangnam-ku, Seoul, 135-080 Republic of Korea

I/We, the undersigned, do hereby appoint the above-identified agent (common representative) as my/our agent/common representative to act for me/us in proceedings concerning all of my/our International Application set forth below.

International Application No.: PCT/KR00/00664

Title of Invention: DECORATIVE FLOOR COVERING COMPRISING

POLYETHYLENE TEREPHTHALATE FILM LAYER IN SURFACE LAYER AND METHOD FOR

PREPARING THE SAME

This 12th day of Nov., 2001

Applicant Name: LG CHEMICAL CO., LTD.

Representative: NO, Ki-Ho

Address: LG Twin Tower, Yoido-dong 20, Youngdungpo-ku, Seoul,

150-721, Republic of Korea

### PCT

### INFORMATION CONCERNING ELECTED OFFICES NOTIFIED OF THEIR ELECTION

(PCT Rule 61.3)

#### From the INTERNATIONAL BUREAU

To

KIM, Won-Ho 7th Fl., Teheran Bd. 825-33 Yoksam-dong, Kangn Seoul 135-080 RÉPUBLIQUE DE CORÉE



Date of mailing (day/month/year)

22 February 2001 (22.02.01)

Applicant's or agent's file reference

OPP000458KR

International application No.

PCT/KR00/00664

International filing date (day/month/year)
26 June 2000 (26.06.00)

Priority date (day/month/year)
26 June 1999 (26.06.99)

IMPORTANT INFORMATION

**Applicant** 

LG CHEMICAL LTD. et al

1. The applicant is hereby informed that the International Bureau has, according to Article 31(7), notified each of the following Offices of its election:

EP:AT,BE,CH,CY,DE,DK,ES,FI,FR,GB,GR,IE,IT,LU,MC,NL,PT,SE National:CN,JP,US

2. The following Offices have waived the requirement for the notification of their election; the notification will be sent to them by the International Bureau only upon their request:

National:BR

3. The applicant is reminded that he must enter the "national phase" before the expiration of 30 months from the priority date before each of the Offices listed above. This must be done by paying the national fee(s) and furnishing, if prescribed, a translation of the international application (Article 39(1)(a)), as well as, where applicable, by furnishing a translation of any annexes of the international preliminary examination report (Article 36(3)(b) and Rule 74.1).

Some offices have fixed time limits expiring later than the above-mentioned time limit. For detailed information about the applicable time limits and the acts to be performed upon entry into the national phase before a particular Office, see Volume II of the PCT Applicant's Guide.

The entry into the European regional phase is postponed until 31 months from the priority date for all States designated for the purposes of obtaining a European patent.

The International Bureau f WIPO 34, chemin des Colombettes 1211 Geneva 20, Switzerland Authorized officer:

Zakaria EL KHODARY

///

Facsimile No. (41-22) 740.14.35

Telephone No. (41-22) 338.83.38

PATENT COOPERATION TREAT

From the

INTERNATIONAL PRELIMINARY EXAMINING AUTHORITY

To:

KIM, Won Ho

7th Fl., Teheran Bldg., 825-33 Yoksam-dong, Kangnam-ku, Seoul 135-080, Republic of Korea

NOTIFICATION OF TRAN INTERNATIONAL PRELIM **EXAMINATION REPOR** 

(PCT Rule 71.1)

Date of mailing

(day/month/year) 28 SEPTEMBER 2001 (28.09.2001)

Applicant's or agent's file reference

OPP000458KR

IMPORTANT NOTIFICATION

International application No. PCT/KR00/00664 International filing date (day/month/year)

Priority date (day/months/year)

26 JUNE 2000 (26.06.2000)

26 JUNE 1999 (26.06.1999)

Applicant

LG CHEMICAL LTD. et al

- The applicant is hereby notified that International Preliminary Examining Authority transmits here with the international preliminary examination report and its annexes, if any, established on the international application.
- A copy of the report and its annexes, if any, is being transmitted to the International Bureau for communication to all the elected Offices.
- Where required by any of the elected Offices, the International Bureau will prepare an English translation of the report(but not of any annexes) and will transmit such translation to those Offices.

#### REMINDER

The applicant must enter the national phase before each elected office by performing certain acts (filing translations and paying national fees) within 30 months from the priority date (or later in some Offices) (Article 39(1)) (see also the reminder sent by the International Bureau with Form PCT/IB/301).

Where a translation of the international application must be furnished to an elected Office, that translation must contain a translation of any annexes to the international preliminary examination report. It is the applicant's responsibility to prepare and furnish such translation directly to each elected Office concerned.

For further details in the applicable time limits and requirements of the elected Offices, see Volume II of the PCT Applicant's Guide.

Name and mailing address of the IPEA/KR

Korean Intellectual Property Office

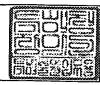
Government Complex-Daejeon, Dunsan-dong, Seo-gu, Daejeon Metropolitan City 302-701, Republic of Korea

Facsimile No. 82-42-472-7140

Authorized officer

COMMISSIONER

Telephone No. 82-42-481-5210



## PATENT COOPERATION TREATY

### **PCT**

### INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Artcle 36 and Rule 70)

[	·			
Applicant's or agent's file reference OPP000458KR		oofTransmittalofInternatio		
International application No. PCT/KR00/00664	International filing date(day/ 26 JUNE 2000 (26.06.2000)		Priority date (day/month/ 26 JUNE 1999 (26.06.1)	•
International Patent Classification (IPC)	or national classification and	IPC	····	<del></del>
IPC7 B32B 27/18				
Applicant				
LG CHEMICAL LTD. et al		···		
This international preliminary examples and is transmitted to the applican	camination report has been part according to Article 36.	repared by this Inter	rnational Preliminary Exa	amining Authority
2. This REPORT consists of a total	of 3 sheets, in	ncluding this cover sl	heet.	
amended and are the basis f 70.16 and Section 607 of the	nied by ANNEXES, i.e., sheet or this report and/or sheets of a Administrative Instructions	ontaining rectification		
These annexes consist of a total of	of sheets.			
IV Lack of unity of inve  V X Reasoned statement citations and explana VI Certain documents of  VII Certain defects in the	f opinion with regard to novel ntion under Article 35(2) with rega tions supporting such stateme	ity, inventive step and ard to novelty, invent ent		licability;
Date of submission of the demand	Dat	e of completion of th	is report	
16 JANUARY 2001 (16.01.2001	)	12 SEPTEMBE	ER 2001 (12.09.2001)	
Name and mailing address of the IPEA/k Korean Intellectual Property Office Government Complex-Daejeon, Dunsan	n-dong, Seo-gu, Daejcon	horized officer  LEE, Jung Hee		APIRA
Metropolitan City 302-701, Republic of Eacsimile No. 82-42-472-7140	· ·	enhone No 82-42-4	81.557 <i>4</i>	

### INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International aplication No.

PCT/KR00/00664

I.	Basi	sis of the report		
1.	With	th regard to the elements of the international application:*		
	X	the international application as originally filed		
	$\overline{\sqcap}$	the description:		
		pages		, as originally filed
		pages,	, filed with the letter of	, filed with the demand
	_		, filed with the fetter of	
		the claims: pages		, as originally filed
		pages	, as amended (together with any s	
		pages		, , filed with the demand
		pages,	, filed with the letter of	
		the drawings:		
		pages		, as originally filed , filed with the demand
		pages,		, fired with the definition
		the sequence listing part of the description:		
		pages		, as originally filed
		pages,	filed with the letter of	, filed with the demand
		pages,	ined with the letter of	
2.	the	ith regard to the language, all the elements marked above were a e international application was filed, unless otherwise indicated mese elements were available or furnished to this Authority in the language of a translation furnished for the purposes of international application (unless than the language of publication of the international application (unless than the language of publication of the international application (unless than the language of publication of the international application).	under this item. te following language <u>English</u> ternational search (under Rule 23.1(b)	which is
		the language of the translation furnished for the purposes of or 55.3).	international preliminary examination	on(under Kules 55.2 ang/
3.		Vith regard to any nucleotide and/or amino acid sequence dis reliminary examination was carried out on the basis of the seq contained in the international application in written form.  filed together with the international application in computer of furnished subsequently to this Authority in written form.	quence listing:	on, the international
			6	
		furnished subsequently to this Authority in computer readable		to discount of the Africa
		The statement that the subsequently furnished written so international applicationas as filed has been furinshed.	equence listing does not go beyond	I the disc losure in the
		The statement that the information recorded in computer r been furnished.	readable form is identical to the writte	en sequence listing has
4.		The amendments have resulted in the cancellation of:		
		the description, pages		
		the claims, Nos.		
		the drawings, sheet		
5.		This opinion has been drawn as if (some of) the amendmen beyond the disclosure as filed, as indicated in the Supplemental Control of the Computer of the Compu		ve been considered to go
*	in thi	placement sheets which have been furnished to the receiving Offic this opinion as "originally filed." and are not annexed to this re I 70.17).		
**	Any i	v replacement sheet containing such amendments must be referre	ed to under item I and annexed to this	report.

#### INTERNATIONAL PRELIMINARY EXAMINATION

International aplication No.

PCT/KR00/00664

V.	Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability;
	citations and explanations supporting such statement

1.	Statement				
	Novelty (N)	Claims	1-31	Y	ES
		Claims		NO	Э
	Inventive step (IS)	Claims Claims	1-31	Y	ES
	Industrial applicability (IA)	Claims	1-31		ES
		Claims		No	Э

2. Citations and explanations (Rule 70.7)

D1: JP-A-08-174783 D2: KR-A-97-33802 D3: KR-A-94-13816

Claims 1-31 meet the criteria set out in PCT Article 33(2)-(4), because the prior arts(D1 to D3) do not teach or fairly suggest a decorative floor covering comprising a surface layer and a substrate layer underneath, wherein the surface layer comprises a polyethylene terephthalate film layer, and a method of preparing the same, and any of the embodiments as specifically set forth in the claims.



### INTERNATIONAL SEARCH REPORT

(PCT Article 18 and Rules 43 and 44)

Applicant's or agent's file reference		of Transmittal of International Search Report					
OPP000458KR	11011011	1/220) as well as, where applicable, item 5 below.					
International application No.	International filing date (day/month/year)	(Earliest) Priority Date (day/month/year)					
PCT/KR00/00664	26 JUNE 2000 (26.06.2000)	26 JUNE 1999 (26.06.1999)					
Applicant							
LG CHEMICAL LTD. et al							
This International search report has been pre to Article 18. A copy is being transmitted to		nority and is transmitted to the applicant according					
This international search report consists of a	<del></del>						
It is also accompanied by a co	py of each prior art document cited in this	report.					
language in which it was filed, unle the international search was ca Authority (Rule 23.1(b)).	ess otherwise indicated under this item. rried out on the basis of a translation of the	basis of the international application in the international application furnished to this					
b. With regard to any nucleotide and was carried out on the basis of the		international application, the international search					
contained in the international	application in written form.						
filed together with the internal	tional application in computer readable for	m.					
furnished subsequently to this	Authority in written form.						
furnished subsequently to this	Authority in computer readable form.						
the statement that the subseque international application as file	enlty furnished written sequence listing de ed has been furnished.	oes not go beyond the disclosure in the					
the statement that the informa furnished.	tion recorded in computer readable form is	identical to the written sequence listing has been					
2. Certain claims were found u	insearchable (See Box I).						
3. Unity of invention is lacking	(See Box II).						
4. With regard to the title,							
X the text is approved as submit	led by the applicant.						
the text has been established b	by this Authority to read as follows:						
5. With regard to the abstract,							
X the text is approved as submit							
[		y as it appears in Box III. The applicant may,					
within one month from the dat	te of mailing of this international search re	port, submit comments to this Authority.					
6. The figure of the drawing to be pub	lished with the abstract is Figure No	2					
X as suggested by the applicant.		None of the figures.					
because the applicant failed to	suggest a figure.	_					
because this figure better char	acterizes the invention.						

International application No. PCT/KR00/00664

		PCT/KR0	0/00664				
A. CLASSIFICATION OF SUBJECT MATTER							
IPC7 B32B 27/18							
According to International Patent Classification (IPC) or to both national classification and IPC							
B. FIELDS SEARCHED							
Minimun documentation searched (classification system followed by classification symbols)							
IPC7 B32B 2	27/08, B32B 27/00, B32B 33/00						
Documentatio	n searched other than minimun documentation to the	extent that such documents are included in	the fileds searched				
	nts and Applications for Invention since 1975 by Models and Applications for Utility Models since 1	975					
Electronic dat NPS, PAJ	a base consulted during the intertnational search (name	ne of data base and, where practicable, sea	rch trerms used)				
C. DOCUI	MENTS CONSIDERED TO BE RELEVANT						
Category*	Citation of document, with indication, where ap	propriate, of the relevant passages	Relevant to claim No.				
Α	JP 08-174783 A (Dainippon Printing Co., Ltd.) See Claim1-9, Fig.1-2		1				
Α	KR 97-33802 A (Hyundai Motor Co., Ltd.) See Abatract, Claim1-3		1				
Α	KR 94-13816 A (Lucky Co., Ltd.) See Abatract, Claim1-10, Fig.1-4		1				
Further	documents are listed in the continuation of Box C.	See patent family annex.					
* Special cat "A" document of to be of par "E" earlier app filing date "L" document of cited to est special rea "O" document means "P" document p	egories of cited documents: lefining the general state of the art which is not considered ticular relevence lication or patent but published on or after the international which may throw doubts on priority claim(s) or which is ablish the publication date of citation or other son (as specified) referring to an oral disclosure, use, exhibition or other bublished prior to the international filing date but later ority date claimed	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevence; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevence; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art "&" document member of the same patent family					
Date of the actual completion of the international search  Date of mailing of the international search report							
09 OCTOBER 2000 (09.10.2000)		10 OCTOBER 2000 (10.10.1	2000)				
	lling address of the ISA/KR	Authorized officer	All mirror				
Government (	trial Property Office Complex-Taejon, Dunsan-dong, So-ku, Taejon City 302-701, Republic of Korea	LEE, Jung Hee	(ORB)				

Telephone No. 82-42-481-5574

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Information on patent family members

International application No. PCT/KR00/00664

	Patent document cited in search report	Publication date	Patent family member(s)	Publication date
	JP 08-174783	09.07.96	None	
	KR 97-33802	22.07.97	None	
	KR 94-13816	16.07.94	None	
- 1				



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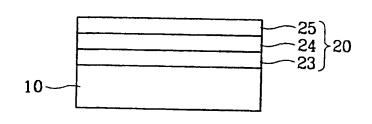
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(54) Title: DECORATIVE FLOOR COVERING COMPRISING POLYETHYLENE TEREPHTHALATE FILM LAYER IN SUR-FACE LAYER AND MANUFACTURING METHOD OF THE SAME



(57) Abstract: The present invention provides a decorative floor covering comprising a surface layer and a substrate layer underneath, wherein the surface layer comprises a polyethylene terephthalate film layer, and a the present invention provides a decorative floor covering comprising a surface layer and a substrate layer underneath, wherein a light weight back layer is further comprised under the substrate layer, and a method for preparing the same. A decorative floor covering of the present invention is a decorative floor covering which has superior surface heat resistance, printing sharpness and realism, and it can be installed conveniently since not only can non-foamed polyvinyl chloride resin be used as a substrate layer, but also the weight of a floor covering is light when a light weight back layer is comprised under the method for preparing the same. Furthermore,

used as a substrate layer, but also the weight of a floor covering is light when a light weight back layer is comprised under the substrate layer.



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(74) Agent: KIM, Won-Ho; 7th Fl., Teheran Bd., 825-33 Yoksam-dong, Kangnam-ku, Seoul 135-080 (KR).

(81) Designated States (national): BR, CN, JP, US.

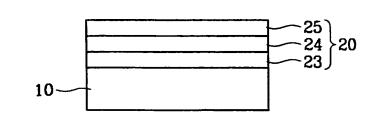
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(54) Title: DECORATIVE FLOOR COVERING COMPRISING POLYETHYLENE TEREPHTHALATE FILM LAYER IN SURFACE LAYER AND MANUFACTURING METHOD OF THE SAME



(57) Abstract: The present invention provides a decorative floor covering comprising a surface layer and a substrate layer underneath, wherein the surface layer comprises a polyethylene terephthalate film layer, and a method for preparing the same. Furthermore, the present invention provides a decorative floor covering comprising a surface layer and a substrate layer underneath, wherein a light weight back layer is further comprised under the substrate layer, and a method for preparing the same. A decorative floor covering of the present invention is a decorative floor covering which has superior surface heat resistance,

printing sharpness and realism, and it can be installed conveniently since not only can non-foamed polyvinyl chloride resin be used as a substrate layer, but also the weight of a floor covering is light when a light weight back layer is comprised under the substrate layer.



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# DECORATIVE FLOOR COVERING COMPRISING POLYETHYLENE TEREPHTHALATE FILM LAYER IN SURFACE LAYER AND MANUFACTURING METHOD OF THE SAME

#### 5 CROSS REFERENCE TO RELATED APPLICATION

This application is based on Korean patent application Nos. 10-1999-0024426 and 10-2000-0034772 filed on June 26, 1999 and June 23, 2000 respectively, which are incorporated hereinto by reference.

#### **BACKGROUND OF THE INVENTION**

#### (a) Field of the Invention

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The present invention relates to a decorative floor covering, more particularly to a decorative floor covering comprising a polyethylene terephthalate film layer in a surface layer, and a manufacturing method of the same.

#### (b) Description of the Related Art

As demands for convenience and health functions in home environments increase in conjunction with recent income level improvements, the need for light weight decorative floor coverings and various functional products used in a common residence or commercial space tends to be increasing.

A decorative floor covering which is installed in common residences and a semi-commercial spaces provides simple functionalities such as durability, stain resistance and fashion characteristics.

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Methods for printing decorative patterns in conventional decorative floor coverings include directly printing patterns on a polyvinyl chloride resin sheet, directly printing patterns on a substrate layer comprising glass fiber etc., and transferring printed patterns on paper or polyester film to a polyvinyl chloride resin sheet surface. However, there have been problems in that these methods do not satisfy human desires for natural patterns without leaving artificial traces. Particularly, although effects of a pattern which is printed on paper or polyester transfer paper is superior, there have been problems in that realism decreases since pattern changes occur after transfer. Furthermore, although a polyvinyl chloride resin layer is formed by adding fillers to a polyvinyl chloride resin sheet layer to raise the transferring effects since complete transfer does not occur due to transferring surface irregularities, the transferred patterns deteriorate due to the fillers.

Secondly, conventional decorative floor coverings in which an olefin resin skin layer such as a transparent polyvinyl chloride resin, polyethylene, etc. is used on a printing layer so as to protect the printing layer tend to generate microscopic air pockets during processing, so a yellowing phenomenon occurs due to the heat changes during processing. In addition, a certain thickness should be maintained in order to provide durability during use, resulting in the problem that transparency is further deteriorated due to the thickness.

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Although there have been cases that non-foaming polyvinyl chloride resin or other materials have been used as a skin layer to improve heat resistance, problems of heat resistance have not been fundamentally solved.

Furthermore, a non-foaming polyvinyl chloride resin layer has been 5 used under a conventional decorative floor covering as a balance layer, and there have been installing problems since the product weight is increased due to the resin layer.

FIG. 1 is a cross-sectional view of a conventional decorative floor covering, wherein a substrate impregnated layer 1 is positioned in the middle. 10 a non-foaming chip layer 21 having a pastel tone and decorative patterns, a durability provided polyvinyl chloride resin skin layer 22, and a durability and heat resistance provided surface treated layer 25 are positioned in order on the substrate impregnated layer 1, and a balance layer 31 which maintains product balance is positioned under the substrate impregnated layer 1.

15

The above conventional decorative floor coverings are manufactured by a method comprising the steps of heat pressing and gelling after fully impregnating glass fiber, pulp, etc. having large pores into a polyvinyl chloride resin sol to make a substrate impregnated layer 1, forming polyvinyl chloride resin chips on non-foamed chips 21 by applying heat after coating 20 calcium carbonate (CaCO<sub>3</sub>) contained polyvinyl chloride resin chips having 5 to 7 colors to a thickness of 1.0 to 1.5 mm on the substrate impregnated layer 1 using a rotary screen roll, laying up a transparent polyvinyl chloride resin sheet on the non-foamed chips 21 as a skin layer 22, laying up a calcium carbonate filled polyvinyl chloride resin sheet under the substrate

impregnated layer 1 as a balance layer 31, and forming a surface treated layer 25 by curing after coating urethane acryl resin, etc. on the skin layer 22.

The conventional decorative floor coverings having these structures have problems in that realism is decreased in the decorative patterns of the non-foamed chip layer 21, transparency of the durability providing polyvinyl chloride resin skin layer 22 is decreased, and its heat resistance is insufficient.

#### **SUMMARY OF THE INVENTION**

It is an object of the present invention to provide a decorative floor covering having a superior heat resistant surface and a method for manufacturing the same, considering problems of the conventional technologies.

It is another object of the present invention to provide a decorative floor covering having superior surface printing sharpness and realism, and a method for manufacturing the same.

It is another object of the present invention to provide a light weight decorative floor covering which can be installed conveniently since non-foamed polyvinyl chloride resin is used as a substrate layer, and a method for manufacturing the same.

In order to accomplish the above objects, the present invention provides a decorative floor covering comprising a surface layer and a substrate layer, wherein the surface layer comprises a polyethylene terephthalate film layer.

Furthermore, the present invention provides a decorative floor

covering comprising a surface layer and a substrate layer, wherein a light back layer is further comprised underneath the substrate layer.

#### BRIEF DESCRIPTION OF THE DRAWINGS

- A more complete appreciation of the invention, and many of the attendant advantages thereof, will be readily apparent as the same becomes better understood by reference to the following detailed description when considered in conjunction with the accompanying drawings, wherein:
- FIG. 1 is a cross-sectional view of a conventional decorative floor 10 covering;
  - FIG. 2 is a cross-sectional view of the decorative floor covering of EXAMPLE 1;
- FIG. 3 is a cross-sectional view of a decorative floor covering in which a glass fiber scrim or a woven or non-woven glass fiber interposed in the middle of a polyvinyl chloride resin substrate layer of the present invention;
  - FIG. 4 is a cross-sectional view of the decorative floor covering of EXAMPLE 2;
- FIG. 5 is a cross-sectional view of the decorative floor covering of 20 EXAMPLE 3; and
  - FIG. 6 is a cross-sectional view of the decorative floor covering of EXAMPLE 4.

#### **DETAILED DESCRIPTION OF THE INVENTION**

In the following detailed description, only the preferred embodiments

of the invention have been shown and described, simply by way of illustration of the best mode contemplated by the inventor(s) of carrying out the invention. As will be realized, the invention is capable of modification in various obvious respects, all without departing from the invention.

5 Accordingly, the description is to be regarded as illustrative in nature, and not restrictive.

The present invention is described in detail as following.

The present invention is a decorative floor covering downwardly comprising a surface layer 20 with a polyethylene terephthalate film having superior transparency, heat resistance, durability, and chemical resistance, and a polyvinyl chloride substrate layer 10 having a high filler content. That is, it is an aim that at the point when a high temperature is applied to the surface of conventional polymer plastic floor coverings, the high temperature is transferred so that not only do carbonization phenomena not occur, but the heat is transferred very quickly. So, a UV coated polyethylene terephthalate film having superior heat resistance is used in the surface layer 20, and polyvinyl chloride resin having a high filler content is used as a substrate so as to swiftly transfer a high temperature applied to the floor covering surface so that the decorative floor covering surface is not carbonized even at a high temperature. Furthermore, patterns of a decorative floor covering are vividly expressed using superior transparency and printability of a polyethylene terephthalate film.

Therefore, printing sharpness and realism are high and heat resistance is improved in a decorative floor covering of the present invention.

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The present invention is described in detail with drawings as follows.

FIG. 2 is a cross-sectional view illustrating one embodiment of the present invention, wherein a polyvinyl chloride resin substrate layer 10 is disposed in a decorative floor covering, and a surface layer 20 is disposed on the substrate layer 10. The surface layer 20 can be further subdivided. That is, the surface layer 20 comprises a surface treated layer 25 on a polyethylene terephthalate film layer 24, and a polyvinyl chloride resin intermediate layer 23 under that.

A method for manufacturing a decorative floor covering of FIG. 2

10 basically comprises the steps of:

- a) manufacturing a polyvinyl chloride resin substrate layer 10;
- b) manufacturing a pressed sheet by laying up a polyvinyl chloride resin intermediate layer 23 on the substrate layer 10, applying pressure, and pressing;
- c) manufacturing a half-finished sheet by laying up a polyethylene terephthalate film layer 24 under which a certain pattern is printed on the pressed sheet of step b), applying pressure, and pressing; and

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d) forming a surface treated layer 25 by UV curing after coating a surface treated layer 25 composition on the half-finished sheet of step c).

In the pressing of step b) and step c), the pressing objects are preferably preheated to a temperature of 140 to 170 °C before laying up, and the applied pressure is preferably from 3 to 10 kgf/cui² during pressing.

The polyvinyl chloride resin substrate layer 10 fulfils its function to dissipate heat applied to the top of the surface layer 20, and it is manufactured by adding a high amount of fillers. Usable fillers include one or more inorganic fillers selected from the group consisting of calcium 5 carbonate, talc, wollastonite, and silica, one or more metallic powders selected from the group consisting of aluminum, copper, and iron, and a mixture thereof. In the fillers used, an inorganic filler is preferably from 50 to 400 weight parts based on 100 weight parts of polyvinyl chloride resin, and a metallic powder is preferably from 5 to 30 weight parts based on 100 weight parts of polyvinyl chloride resin. In order to improve processabilities of a calender, the consumed amount can be increased 2 to 5 times more by processing after premixing a metallic powder, a resin, and a plasticizer than when a metallic powder alone is used.

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A preferable method embodiment for manufacturing a polyvinyl 15 chloride resin substrate layer 10 is manufacturing a sheet by adding a plasticizer for reinforcing product flexibility, 30 to 50 weight parts of dioctyl phthalate, heat resistant stabilizers for providing heat resistant stabilities, 3 to 5 weight parts of a barium-zinc based compound and 3 to 5 weight parts of epoxy compound, a filler for providing product hardness and heat resistant 20 dispersibility, 50 to 400 weight parts of calcium carbonate (CaCO<sub>3</sub>), and 3 to 5 weight parts of pigment for imparting surface colors, to 100 weight parts of polyvinyl chloride resin, fully kneading at a rolling mill at 160 to 190  $\,^\circ\mathrm{C}\,$ , and rolling to a thickness of 0.8 to 1.3 mm.

Preferably 50 to 400 weight parts of a calcium carbonate filler are

used based on 100 weight parts of polyvinyl chloride resin since the higher the content of calcium carbonate, not only does the heat conductibility increase, but also press marks caused by heavy objects such as furniture used in everyday life are minimized due to an increased surface hardness. 5 However, when 400 or more weight parts of calcium carbonate are used based on 100 weight parts of polyvinyl chloride resin, the polyvinyl chloride resin does not chemically bond with the calcium carbonate, weakening the cohesion of the polyvinyl chloride resin resulting in a processibility decrease, even though it is profitable in the aspect of manufacturing cost.

Furthermore, glass fiber scrim, or woven or nonwoven long glass fiber11 can be interposed in the middle of the polyvinyl chloride resin substrate layer 10 for dimensional stability, wherein the interposition method is pressing and applying pressure of 3 to 10 kgf/cm² at an embossing roll under the state of latent heat of 100 to 150 ℃ after rolling the first polyvinyl 15 chloride resin sheet at a calender. Thereafter, the second polyvinyl chloride resin sheet is pressed under a glass scrim layer, or a woven or nonwoven glass fiber 11, thus interposing it inside a polyvinyl chloride resin substrate layer 10, and the second polyvinyl chloride resin sheet under which a separate light back layer 30, usually a fiber layer 35, can be attached in 20 advance.

A structure of a decorative floor covering of the present invention in which a glass fiber scrim or woven or nonwoven glass fiber 11 is interposed in a polyvinyl chloride resin substrate layer 10 is illustrated in FIG. 3 and FIG.

4.

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The polyvinyl chloride resin intermediate layer 23, which is used on a polyvinyl chloride resin substrate layer 10 so that color of a substrate layer 10 can be concealed in order to support printed pattern sharpness and realism of an upper surface layer 20 has the additional function of dissipating heat when heat is applied to the upper surface layer 20, and it also transfers heat to the polyvinyl chloride resin substrate layer 10 positioned underneath.

For this, a 0.1 to 1 mm thick sheet is manufactured by rolling a polyvinyl chloride resin composition comprising 100 weight parts of polyvinyl chloride resin, 25 to 50 weight parts of dioctyl phthalate, 50 to 150 weight 10 parts of calcium carbonate, 3 to 5 weight parts of titanium oxide, and 2 to 5 weight parts of heat stabilizer in a calender.

More particularly, after adding 25 to 50 weight parts of dioctyl phthalate plasticizer for reinforcing product flexibility, 2 to 5 weight parts of barium-zinc (Ba-Zn) based stabilizer and 2 to 5 weight parts of epoxy stabilizer heat resistant stabilizers for providing heat resistant stability, 50 to 150 weight parts of inorganic filler such as calcium carbonate etc. for providing product hardness and heat resistant dispersibility, and 3 to 5 weight parts of titanium oxide (TiO<sub>2</sub>) pigment for making the surface color white, to a main raw material of polyvinyl chloride and sufficiently kneading at a rolling 20 roll at 160 to 190 °C, a polyvinyl chloride resin intermediate layer 23 is manufactured by rolling to 0.1 to 1 mm, more preferably to a 0.1 to 0.2 mm thickness in a calender.

The polyethylene terephthalate film layer 24, under which highly realistic and vivid decorative patterns are printed by a printing method such

as gravure printing method, etc., is positioned on the polyvinyl chloride resin intermediate layer 33.

It is preferable to use this polyethylene terephthalate film layer 24 after coating a primer selected from the group consisting of polyvinyl acetate 5 based, polyurethane acrylate based, and ethylenevinyl acetate based primers to a coating thickness of 0.1 to 10  $\,\mu\mathrm{m}$  for adhesion to an UV surface treated layer 25 positioned at the upper surface of the film layer and to a polyvinyl chloride resin intermediate layer 23 positioned under the film layer.

Printing certain patterns on a polyethylene terephthalate film layer 24 10 is possible before or after the primer treatment, and the polyethylene terephthalate film layer is used during pressing by winding on a paper pipe. steel pipe, etc. after manufacturing. Furthermore, printing can be done on the polyvinyl chloride resin intermediate layer 23 as opposed to the polyethylene terephthalate film layer 24.

A thickness of a polyethylene terephthalate film layer 24 is preferably from 10 to 100  $\mu$ m, because when the polyethylene terephthalate film is thinner than 10  $\mu$ m the printing surface is expanded by tension during layingup, and when it is thicker than 100  $\mu m$  product surface folding occurs when the product is folded since polyethylene terephthalate film has low flexibility 20 when it is thicker than 100  $\mu$ m.

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The surface treated layer 25 which is used to provide stain resistance or durability synergistic effectsis formed by coating an UV surface treated layer 25 composition on the polyethylene terephthalate film layer 24, and

curing with ultraviolet rays, etc. A UV surface treated layer 25 composition is preferably selected from the group consisting of urethane acrylate, silicone acrylate, and epoxy acrylate, and preferably further comprising acryl based or urethane based beads having a particle size of 5 to 20  $\mu$ m, in order to maintain the heat resistant function. It is cross linking cured with ultraviolet rays or electron beams: urethane acryl resin is cured by ultraviolet ray irradiation depending on composition, and aqueous urethane resin can be cured with heat.

Ease of installation can be provided by additionally using a cork

10 balance layer 33 as a light back layer 30, or a wooden powder balance layer

34, or a fabric layer 35 underneath the polyvinyl chloride resin substrate layer

10, thus reducing the weight of the decorative floor covering while

maintaining the same total thickness. The light back layer 30 consisting of a

cork balance layer 33, wooden powder balance layer 34, or a fabric layer 35

15 can provide anti-insect effects, heat insulating effects, a humidity controlling

function, and impact absorbing effects as well.

For this, the cork balance layer 33 is manufactured in a sheet which is cut to a thickness of 1.0 to 2.0 mm using a slicing machine after a cork layer of a natural tree is peeled off, and pulverized to a particle size of 5 to 10 meshes, put into a large cylinder together with adhesive, and bonded by applying pressure. A cork having preferably 2% or less moisture content is used since a higher moisture content causes higher contraction or expansion due to heat. A cork balance layer 33 can be manufactured to a thickness of 3.0 to 4.0 mm so as to further improve performance aspects such as sound

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insulation, impact absorption, etc. This cork balance layer 33 can improve the installation property of product weight since its specific gravity is over 8 to 10 times less than a polyvinyl chloride resin sheet containing general inorganic materials.

Furthermore, the wooden powder balance layer 34 which maintains the product balance and provides anti-insect effects, heat insulating effects, a humidity controlling function, impact absorbing effects, etc. is manufactured in a sheet which is rolled to a thickness of 0.8 to 1.0 mm using a two sets of pressure rolls after adding 100 to 150 weight parts of wooden powder having 10 a particle size of 200 to 300 meshes, 3 to 5 weight parts of heat resistant barium-zinc based compound as a stabilizer, 1 to 3 weight parts of internal antiadditive stearic acid, 30 to 50 weight parts of plasticizer dioctyl phthalate, and 3 to 5 weight parts of other pigment to 100 weight parts of polyvinyl chloride resin, and kneading it together, wherein the wooden powder is 15 coating treated to prevent moisture absorption with 30 to 50 weight% of surfactant, based on the wooden powder weight. Polyethylene glycol, polybutyl glycol, polymethyl glycol, polymethyl propylene glycol, etc. can be used as a surfactant. Furthermore, surfactants having preferably 2% or less moisture content are used since severe contraction or expansion by heat can 20 occur when moisture content is higher. The wooden powder balance layer 34 can be manufactured to a thickness of 3,0 to 4.0 mm to further improve functionality aspects such as sound insulation, impact absorption, etc. This wooden powder balance layer 34 can improve the installation property of product weight since its specific gravity is over 8 to 10 times less than a

polyvinyl chloride resin sheet containing general inorganic materials.

Furthermore, the fiber layer 35 which prevents product deformation by dimensional change and provides the product with lightness uses woven or nonwoven polyester, woven or nonwoven polypropylene, woven or nonwoven glass fiber, etc. This fabric layer 35 is used by fixing it with an adhesive after coating a plastisol such as polyvinyl chloride, etc. under the polyvinyl chloride resin substrate layer 10, and simultaneously gelling the sol and pressing with the fiber layer 35, or preparing it under the cork balance layer 33 or the wooden powder balance layer 34. The woven thickness of the fiber layer 35 is preferably from 10s x 10s to 25s x 15s, and the density is preferably from 20 x 20 to 30 x 30 roll/inch.

This light back layer 30 is adhered under the polyvinyl chloride resin substrate layer 10 using an adhesive. The adhesive layer 32 is preferably a sheet or impregnated sheet which is rolled to a thickness of 0.03 to 0.2 mm

15 by a 4 roll calender roll or an extruder at a temperature of 80 to 160 °C using urethane resin, acryl resin, vinyl acetate resin, etc. considering the melting point of each resin, and a light back layer 30 can be adhered by separately coating a plastisol such as polyvinyl chloride, etc. under a polyvinyl chloride resin substrate layer 10 and using this gelled sol. In case of using a rolled sheet or an impregnated sheet as an adhesive layer 32, a pressed product or half-finished product to be adhered is preheated, an adhesive layer 32 is positioned under a substrate layer 10, laid up, and pressed with a pressure of 3 to 10 kg/cm<sub>2</sub>.

A decorative floor covering of the present invention is manufactured

by the various manufacturing methods depending on the application method of the light back layer 30. These manufacturing methods can be represented in the following various embodiments in addition to the above described basic manufacturing method.

One embodiment of a manufacturing method of a decorative floor covering of the present invention comprises the steps of:

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- a) manufacturing the first polyvinyl chloride resin substrate layer 10;
- b) pressing by applying pressure in a embossing roll after positioning a glass fiber scrim, or a woven or unwoven glass fiber 11 underneath the first substrate layer 10 under the condition that a surface of the first polyvinyl chloride resin substrate layer 10 of step a) maintains latent heat of 100 to 150 °C;
- c) manufacturing the second polyvinyl chloride resin substrate layer
   10;
- d) coating a plastisol underneath the second polyvinyl chloride resin substrate layer 10, positioning a light back layer 30 of a fiber layer 35, applying pressure, and pressing with a press roll at a heating drum at 140 to 150 ℃;
  - e) manufacturing a polyvinyl chloride resin intermediate layer 23;
- f) manufacturing a pressed sheet by applying pressure at a press after downwardly positioning a polyvinyl chloride resin intermediate layer 23, the first polyvinyl chloride resin substrate layer 10 under which glass fiber scrim, or woven or unwoven glass fiber 11 is pressed, the second polyvinyl chloride resin substrate layer 10

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under which a light back layer 30 of a fiber layer 35 is pressed, and preheating to a temperature of 140 to 170  $\,^{\circ}$ C;

- g) manufacturing a half-finished sheet by applying pressure and pressing at an embossing roll after preheating the pressed sheet of step f) to a temperature of 140 to 170 ℃ and positioning a polyethylene terephthalate film layer 24 on a polyvinyl chloride resin intermediate layer 23; and
- h) forming a surface treated layer 25 by UV (ultraviolet ray) curing after coating a surface treated layer 25 composition on the half-finished sheet of step g).

Another embodiment of a manufacturing method of a decorative floor covering of the present invention comprises the steps of:

- a) manufacturing a polyvinyl chloride resin substrate layer 10;
- b) laying up a polyvinyl chloride resin intermediate layer 23 on the preheated substrate layer 10 of step a), applying pressure, and pressing;
- c) manufacturing a half-finished sheet by laying up a polyethylene terephthalate film layer 24 on the preheated polyvinyl chloride resin intermediate layer 23 of step b), applying pressure, and pressing;
- d) laying up an adhesive layer 32 underneath the preheated halffinished sheet of step c), applying pressure, and pressing;
- e) laying up a light back layer 30 underneath the adhesive layer 32 of step d), applying pressure, and pressing; and
- f) forming a surface treated layer 25 after coating a surface treated

layer 25 composition on the pressed polyethylene terephthalate film layer 24 of step e), and curing.

In the manufacturing method, a method for laying up and pressing a light back layer 30 can be selected from a method for laying up and pressing a fabric layer 35 underneath a wooden powder balance layer 34 after laying up and pressing a cork balance layer 33 underneath an adhesive layer 32, or laying up and pressing a wooden powder balance layer 34 underneath an adhesive layer 32.

In the manufacturing methods, the preheating in each step is carried out at a temperature preferably from 140 to 170 °C considering the melting point of polyvinyl chloride resin, and an applying pressure during pressing after preheating is preferably from 3 to 10 kg/cm<sub>2</sub>.

A surface treated layer 25, a polyethylene terephthalate film layer 24, a polyvinyl chloride resin intermediate layer 23, a substrate layer 10, and an adhesive layer 32 in a decorative floor covering of the present invention are manufactured in a thickness preferably from 1 to 1.5 mm.

The present invention is described further in detail through the following EXAMPLES and COMPARATIVE EXAMPLES. However, the following EXAMPLES are only for exemplifying the present invention, and the present invention is not limited to the following EXAMPLES.

#### [EXAMPLES]

#### **EXAMPLE 1**

(Manufacturing of a polyvinyl chloride resin substrate layer 10)

A polyvinyl chloride resin substrate layer 10 sheet having a thickness

of 0.75 mm was manufactured by kneading 100 weight parts of polyvinyl chloride resin having a degree of polymerization of 1000, 42 weight parts of dioctyl phthalate, 250 weight parts of calcium carbonate powder, and 2 weight parts of heat resistant stabilizer in a Banbury mixer, and rolling the softened and molten compound with a calender at 130 to 170 °C.

(Manufacturing of a polyvinyl chloride resin intermediate layer 23)

A polyvinyl chloride resin intermediate layer 23 sheet having a thickness of 0.1 mm was manufactured by kneading 100 weight parts of polyvinyl chloride resin having a degree of polymerization of 1000, 38 weight parts of dioctyl phthalate, 90 weight parts of calcium carbonate powder, 2.5 weight parts of heat resistant stabilizer, and 10 weight parts of titanium oxide in a Banbury mixer, and rolling the softened and molten compound with a calender at 130 to 170 °C.

(Pressing of a polyvinyl chloride resin intermediate layer 23)

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After preheating the manufactured polyvinyl chloride resin substrate layer 10 sheet at a temperature of 150 to 170 °C, a polyvinyl chloride resin intermediate layer 23 was positioned on the preheated substrate layer 10, laid up, and pressed at a pressure of 3 to 6 kg/cm².

(Manufacturing of a polyethylene terephthalate film layer 24)

Certain patterns were imparted in the gravure printing method under a polyethylene terephthalate film (SG88 manufactured by SK Corporation) having a thickness of 50  $\mu$ m on both sides of which an acryl and polyester based primer is treated to an average thickness of 0.5  $\mu$ m.

(Pressing of a polyethylene terephthalate film layer 24)

After preheating a sheet in which a polyvinyl chloride resin intermediate layer 23 was pressed on the manufactured polyvinyl chloride resin substrate layer 10 sheet at a temperature of 150 to 170 °C, the polyethylene terephthalate film layer 24 under which certain patterns were imparted was positioned on the preheated sheet, laid up, and pressed at a pressure of 3 to 6 kg/cui².

(Heat resistant UV (ultraviolet rays) coating treatment)

#### 15 EXAMPLE 2

(Manufacturing of a polyvinyl chloride resin substrate layer 10)

The same polyvinyl chloride resin substrate layer 10 sheet as in the EXAMPLE 1 was manufactured.

(Attaching of woven cloth 30)

After coating a polyvinyl chloride sol (plastisol) under the polyvinyl chloride resin substrate layer 10 sheet, a polyester woven cloth (thickness: 20s x 20s, density: 25 x 25 /inch) was positioned and pressed with a press roll at a heating drum at 140 to 150 °C.

(Pressing of glass fiber scrim 11)

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One polyvinyl chloride resin substrate layer 10 sheet was additionally manufactured by rolling a polyvinyl chloride substrate layer 10 sheet using a calender in the same method as in the EXAMPLE 1, and pressed by applying pressure while passing the sheet and glass fiber scrim (average weight of 200 g/m²) which is positioned underneath the sheet through an embossing roll (pressure of 10 kgf/cm²) in the state of sheet formation, i.e., in the state that a latent sheet surface temperature of 100 to 120 °C is maintained.

(Manufacturing of a polyvinyl chloride resin intermediate layer 23)

The same polyvinyl chloride resin intermediate layer 23 as in the EXAMPLE 1 was manufactured.

Each of a polyvinyl chloride intermediate layer 23, a polyvinyl chloride resin substrate layer 10 sheet under which a glass fiber scrim 11 is pressed, and a polyvinyl chloride resin substrate layer 10 sheet under which polyester woven cloth is pressed were respectively preheated to a temperature of 140 to 170 °C, downwardly positioned in order, laid up, and pressed by passing through an embossing roll at 160 °C (pressure of 10 kgf/cu²).

(Manufacturing of a polyethylene terephthalate film layer 24)

The same polyethylene terephthalate film layer 24 as in the EXAMPLE 1 was prepared.

(Pressing of a polyethylene terephthalate film layer 24)

After preheating the manufactured pressed sheet at a temperature of

150 to 170  $^{\circ}$ C, the polyethylene terephthalate film layer 24 under which certain patterns were imparted was positioned on the preheated sheet, laid up, and pressed at a pressure of 3 to 6 kgf/cu².

(Heat resistant UV coating treatment)

An urethane acrylate based heat resistant UV treating agent in which a heat resistant stabilizer and acryl beads were added and contained was coated on the polyethylene terephthalate film layer 24 pressed sheet to a coating thickness of 25 μm, cured by an UV lamp in the non-oxygen curing method, thus forming a heat resistant UV coating layer 25 on the uppermost layer. The final product thickness was 1.8 mm. This manufactured decorative floor covering is illustrated in FIG. 4.

#### EXAMPLE 3

(Manufacturing of a polyvinyl chloride resin substrate layer 10)

The same polyvinyl chloride resin substrate layer 10 as in the 15 EXAMPLE 1 was manufactured.

(Manufacturing of a polyvinyl chloride resin intermediate layer 23)

The same polyvinyl chloride resin intermediate layer 23 as in EXAMPLE 1 was manufactured.

(Pressing of a polyvinyl chloride resin intermediate layer 23)

A polyvinyl chloride resin intermediate layer 23 sheet was pressed on a substrate layer 10 in the same method as in EXAMPLE 1.

(Manufacturing of a polyethylene terephthalate film layer 24)

The same polyethylene terephthalate film layer 24 as in the EXAMPLE 1 was manufactured.

(Pressing of a polyethylene terephthalate film layer 24)

A half-finished sheet was manufactured by pressing the polyethylene terephthalate film layer 24 on a sheet in which a polyvinyl chloride resin intermediate layer 23 was pressed on the polyvinyl chloride resin substrate

5 layer 10 manufactured in the same method as in EXAMPLE 1.

(Manufacturing an adhesive layer 32)

(Pressing of an adhesive layer 32)

After preheating the polyethylene terephthalate film layer 24 pressed half-finished sheet at a temperature of 150 to 170 °C. the adhesive layer 32 was positioned under the preheated half-finished sheet, laid up, and pressed at a pressure of 3 to 6 kg/cm².

(Manufacturing of a cork balance layer 33)

A sheet which was cut to a thickness of 1 mm with a slicing machine after a cork layer of a natural tree was peeled off, pulverized to a particle size of 5 to 10 meshes, put into a large cylinder together with 5 weight% of vinyl acetate adhesive per cork weight, and bonded by applying pressure.

(Pressing of a cork balance layer 33)

The cork balance layer 33 was positioned underneath a half-finished sheet under which the adhesive layer 32 is pressed, laid up, and pressed at a pressure of 3 to 6 kg/cul².

(Heat resistant UV coating treatment)

An urethane acrylate based heat resistant UV treating agent in which a heat resistant stabilizer and acryl beads were added and contained was coated on the pressed sheet to a coating thickness of 25  $\mu$ m, cured by an UV lamp in the non-oxygen curing method, thus forming a heat resistant UV coating layer 25 on the uppermost layer. The final product thickness was 2.0 mm. This manufactured decorative floor covering is illustrated in FIG. 5.

#### **EXAMPLE 4**

(Manufacturing of a polyvinyl chloride resin substrate layer 10)

The same polyvinyl chloride resin substrate layer 10 as in the 10 EXAMPLE 1 was manufactured.

(Manufacturing of a polyvinyl chloride resin intermediate layer 23)

The same polyvinyl chloride resin intermediate layer 23 as in EXAMPLE 1 was manufactured.

(Pressing of a polyvinyl chloride resin intermediate layer 23)

A polyvinyl chloride resin intermediate layer 23 sheet was pressed on a substrate layer 10 in the same method as in EXAMPLE 1.

(Manufacturing of a polyethylene terephthalate film layer 24)

The same polyethylene terephthalate film layer 24 as in the EXAMPLE 1 was manufactured.

20 (Pressing of a polyethylene terephthalate film layer 24)

A half-finished sheet was manufactured by pressing the polyethylene terephthalate film layer 24 on a sheet in which a polyvinyl chloride resin intermediate layer 23 was pressed on the polyvinyl chloride resin substrate layer 10 manufactured in the same method as in EXAMPLE 1.

(Manufacturing an adhesive layer 32)

A 0.1 mm thick adhesive layer 32 was manufactured by rolling vinylacetate resin using a 4 roll calender at 120  $\,^{\circ}$ C.

(Pressing of an adhesive layer 32)

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After preheating the polyethylene terephthalate film layer 24 pressed half-finished sheet at a temperature of 150 to 170  $^{\circ}$ C, the adhesive layer 32 was positioned under the preheated half-finished sheet, laid up, and pressed at a pressure of 3 to 6 kg/cui.

(Manufacturing of a wooden powder balance layer 34)

10 A wooden powder balance layer 34 sheet was rolled to a thickness of 1 mm using a two sets of pressure rolls at a temperature of 130 to 170 °C after kneading 100 weight parts of polyvinyl chloride resin, 150 weight parts of wooden powder having a particle size of 200 to 300 meshes, 5 weight parts of heat resistant stabilizer barium-zinc based compound, 1 weight part of internal antiadditive stearic acid, 30 weight parts of plasticizer dioctyl phthalate, and 5 weight parts of pigment.

(Pressing of a cork balance layer 33)

The wooden powder balance layer 34 was positioned underneath a half-finished sheet under which the adhesive layer 32 was pressed, laid up, and pressed at a pressure of 3 to 6 kg/cm².

(Manufacturing of fiber layer 35)

A polyvinyl chloride sol (plastisol) coated polyester nonwoven (thickness: 20s x 20s, density: 25 x 25 /inch) was prepared.

(Pressing of a fiber layer 35)

The fiber layer 35 was positioned underneath a half-finished sheet under which the cork balance layer 33 was pressed, laid up, and pressed at a pressure of 3 to 6 kg/cu².

(Heat resistant UV coating treatment)

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An urethane acrylate based heat resistant UV treating agent in which a heat resistant stabilizer and acryl beads were added and contained was coated on the pressed sheet to a coating thickness of 25 µm, cured by an UV lamp in the non-oxygen curing method, thus forming a heat resistant UV coating layer 25 on the uppermost layer. The final product thickness was 2.0 mm. This manufactured decorative floor covering is illustrated in FIG. 6. COMPARATIVE EXAMPLE 1

2.3 mm thick polyvinyl chloride resin decorative cushion floor covering

As a conventional 2.3 mm thick polyvinyl chloride resin decorative cushion floor covering, a Woodleum Plus product manufactured by LG Chemical Ltd. downwardly comprising an UV coating layer, a transparent layer, a printing layer, an upper foaming layer, a substrate layer, an underneath foaming layer, and a sizing layer was prepared.

The Woodleum Plus product was manufactured by the following method.

(Manufacturing of a substrate layer 10)

A substrate layer was manufactured by gelling at a temperature of 150 to 230  $^{\circ}$ C after depositing a sol made by mixing 100 weight parts of

polyvinyl chloride resin having a degree of polymerization of 2000, 10 to 100 weight parts of plasticizer, 1 to 20 weight parts of stabilizer, 1 to 20 weight parts of pigment, 1 to 50 weight parts of calcium carbonate, and other additives on a substrate such as glass fiber, vellum paper, or mineral material paper.

(Manufacturing of an upper foaming layer)

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An upper foaming layer was manufactured by foaming at 170 to 250 ℃ for 30 to 180 seconds after coating a sol made by mixing 100 weight parts of polyvinyl chloride resin having a degree of polymerization of 2000, 10 to 120 weight parts of plasticizer, 1 to 20 weight parts of stabilizer, 1 to 20 weight parts of pigment, 1 to 20 weight parts of foaming agent, and other additives to a thickness of 0.1 to 0.2 mm on the substrate layer.

(Manufacturing of a printing layer and a transparent layer)

A printing layer was formed by printing certain patterns on the upper foaming layer using a gravure or offset ink, a rotary screen, and a transparent layer was manufactured by gelling at 170 to 230 °C for 30 to 180 seconds after coating a sol made by mixing 100 weight parts of polyvinyl chloride resin having degree of polymerization of 2000, 10 to 150 weight parts of plasticizer, 1 to 20 weight parts of stabilizer, and other additives to a thickness of 0.2 to 0.25 mm on the printing layer.

(Pressing of an underneath foaming layer and a sizing layer)

An underneath foaming layer was manufactured by foaming at 170 to 250 ℃ for 30 to 180 seconds after coating a sol made by mixing 100 weight parts of polyvinyl chloride resin having a degree of polymerization of 2000, 10

to 120 weight parts of plasticizer, 1 to 20 weight parts of stabilizer, 1 to 20 weight parts of pigment, 1 to 20 weight parts of foaming agent, and other additives to a thickness of 0.1 to 0.2 mm on the substrate layer, and a sizing layer was manufactured by gelling at 170 to 230 °C for 30 to 180 seconds after coating a sol made by mixing 100 weight parts of polyvinyl chloride resin having degree of polymerization of 2000, 10 to 120 weight parts of plasticizer, 1 to 20 weight parts of pigment, 1 to 20 weight parts of stabilizer, 1 to 100 weight parts of calcium carbonate, and other additives to a thickness of 0.1 to 0.5 mm on the underneath foaming layer.

10 (Manufacturing an UV layer)

A UV layer was manufactured by UV curing unsaturated polyester resin, urethane acryl resin, epoxy acrylate resin, or polyester acryl resin on the transparent layer.

#### COMPARATIVE EXAMPLE 2

A decorative floor covering comprising a 3.0 mm thick polyvinyl chloride resin layer

As a conventional decorative floor covering comprising a 3.0 mm thick polyvinyl chloride resin layer, a Deco Tile manufactured by LG Chemical Ltd. downwardly comprising an UV layer, a transparent film layer, a printing layer, an intermediate layer, a base layer, and a back layer was prepared.

(Manufacturing of a transparent film layer)

A 0.2 to 0.3 mm thick transparent sheet was manufactured by mixing 100 weight parts of polyvinyl chloride resin having a degree of polymerization of 800 to 1000, 10 to 40 weight parts of plasticizer, 1 to 20 weight parts of

stabilizer, and other additives, and sheeting in the continuous calendaring method.

(Manufacturing an UV layer)

An UV layer was manufactured by UV curing unsaturated polyester resin, urethane acryl resin, epoxy acrylate resin, or polyester acryl resin on the transparent layer.

(Structure of other layers)

Other underneath layers were manufactured by an ordinary method, i.e., continuously calendaring non-foaming sheet layers, whereby each layer was manufactured by heat pressing at a temperature of 130 to 160 °C.

The product thickness was 3 mm, and the thickness of a surface skin layer from an uppermost layer, i.e., an UV layer to a transparent film layer was 0.25 mm.

#### **COMPARATIVE EXAMPLE 3**

15 <u>1.8 mm thick polyvinyl chloride resin non foaming decorative floor</u> covering

As a conventional 1.8 mm thick polyvinyl chloride resin non foaming decorative floor covering, a Luckstrong manufactured by LG Chemical Ltd. downwardly comprising an UV layer, a transparent skin layer, a chip layer, a substrate layer, a base sol layer, and a back layer was prepared.

The Luckstrong product was manufactured by the following method.

(Manufacturing of a substrate layer)

A substrate layer was manufactured by gelling at a temperature of 150 to 200 ℃ after depositing a sol made by mixing 100 weight parts of

polyvinyl chloride resin having a degree of polymerization of 2000, 10 to 100 weight parts of plasticizer, 1 to 20 weight parts of stabilizer, 1 to 150 weight parts of calcium carbonate, and other additives on a substrate such as glass fiber, vellum paper, or mineral material paper.

(Manufacturing of a base sol layer and a chip layer)

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After manufacturing a base sol layer by coating a sol made by mixing 100 weight parts of polyvinyl chloride resin having a degree of polymerization of 2000, 10 to 50 weight parts of plasticizer, 1 to 20 weight parts of stabilizer, 1 to 20 weight parts of pigment, 1 to 50 weight parts of calcium carbonate, and other additives to a thickness of 0.1 to 0.2 mm on the substrate layer, a chip layer was manufactured by arranging on the base sol layer chips having a size of 8 x 9 mm which were prepared after manufacturing a 0.8 to 1.2 mm thick transparent sheet by mixing 100 weight parts of polyvinyl chloride resin having a degree of polymerization of 800 to 1000, 10 to 40 weight parts of plasticizer, 1 to 20 weight parts of stabilizer, 50 to 200 weight parts of calcium carbonate, 1 to 5 weight parts of pigment, and other additives, and sheeting in the continuous calendaring method.

(Manufacturing of a transparent skin layer)

A transparent skin layer was manufactured by gelling at 170 to 200 ℃ after coating a sol made by mixing 100 weight parts of polyvinyl chloride resin having a degree of polymerization of 1700 to 2000, 10 to 50 weight parts of plasticizer, 1 to 5 weight parts of stabilizer, and other additives to a thickness of 0.1 to 0.2 mm on the chip layer.

(Manufacturing an UV layer)

An UV layer was manufactured by UV curing unsaturated polyester resin, urethane acryl resin, epoxy acrylate resin, or polyester acryl resin on the transparent layer.

(Pressing of a back layer)

A back layer was manufactured by pressing with heat and pressure after manufacturing a 0.8 to 1.2 mm thick transparent sheet by mixing 100 weight parts of polyvinyl chloride resin having a degree of polymerization of 800 to 1000, 10 to 40 weight parts of plasticizer, 1 to 20 weight parts of stabilizer, 50 to 200 weight parts of calcium carbonate, 1 to 5 weight parts of pigment, and other additives, and sheeting in the continuous calendaring method.

#### **COMPARATIVE EXAMPLE 4**

3.5 mm thick polyvinyl chloride resin decorative cushion floor covering

As a conventional 3.5 mm thick polyvinyl chloride resin decorative cushion floor covering, a Sharpeny product manufactured by LG Chemical Ltd. downwardly comprising an UV layer, a transparent layer, a printing layer, an upper foaming layer, a substrate layer, and a mechanical foaming layer was prepared.

The Sharpeny product was manufactured by the following method.

(Manufacturing of a substrate layer)

A substrate layer was manufactured by gelling at a temperature of 150 to 230 ℃ after depositing a sol made by mixing 100 weight parts of polyvinyl chloride resin having a degree of polymerization of 2000, 10 to 100

weight parts of plasticizer, 1 to 20 weight parts of stabilizer, 1 to 20 weight parts of pigment, 1 to 50 weight parts of calcium carbonate, and other additives on a substrate such as glass fiber, vellum paper, or mineral material paper.

(Manufacturing of an upper foaming layer)

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An upper foaming layer was manufactured by foaming at 170 to 250 °C for 30 to 180 seconds after coating a sol made by mixing 100 weight parts of polyvinyl chloride resin having a degree of polymerization of 2000, 10 to 120 weight parts of plasticizer, 1 to 20 weight parts of stabilizer, 1 to 20 weight parts of pigment, 1 to 20 weight parts of foaming agent, and other additives to a thickness of 1.0 to 2.0 mm on the substrate layer.

(Manufacturing of a printing layer and a transparent layer)

A printing layer was formed by printing certain patterns on the upper foaming layer using a gravure or offset ink, a rotary screen, and a transparent layer was manufactured by gelling at 170 to 230 °C for 30 to 180 seconds after coating a sol made by mixing 100 weight parts of polyvinyl chloride resin having a degree of polymerization of 2000, 10 to 150 weight parts of plasticizer, 1 to 20 weight parts of stabilizer, and other additives to a thickness of 0.2 to 0.25 mm on the printing layer.

20 (Manufacturing of a mechanical foaming layer)

After manufacturing a sol by mixing 100 weight parts of polyvinyl chloride resin having a degree of polymerization of 1100 to 2000, 60 to 80 weight parts of plasticizer, 0 to 30 weight parts of calcium bicarbonate, 2 to 3 weight parts of stabilizer, and other additives, and putting the sol into a

mechanical foaming layer, and manufacturing a cream having a density of 0.3 to 0.7 g/cm² by high speed agitating at 200 to 400 rpm while injecting air at 5 to 8 bar, a mechanical foaming layer was manufactured by coating the cream to a thickness of 2 to 3 mm on the substrate layer, and gelling at 160 to 200 °C.

(Manufacturing an UV layer)

An UV layer was manufactured by UV curing unsaturated polyester resin, urethane acryl resin, epoxy acrylate resin, or polyester acryl resin on the transparent layer.

#### 10 EXAMPLE 5

(Transparency comparison)

In order to confirm vividness of printed patterns and transparency of a polyethylene terephthalate film layer 24 of the present invention, haze values of transparency of the polyethylene terephthalate film layer 24 of the present invention and conventional polyvinyl chloride sheet layer were measured more than 5 times by the method of ASTM D-1003, and their average values are represented in the following Table 1.

A polyethylene terephthalate film (SG88 manufactured by SK Chemical Ltd.) was used in a polyethylene terephthalate film of the present invention, a comparison polyvinyl chloride sheet was manufactured after manufacturing a sol by mixing 100 weight parts of polyvinyl chloride resin having a degree of polymerization of 2000, 10 to 50 weight parts of plasticizer, 1 to 20 weight parts of stabilizer, and other additives, coating the sol to a thickness of 0.2 mm, and gelling in an oven at 200 °C, and the

collected films of the sheet were used as samples.

[Table 1]

Classification	Polyvinyl chloride sheet	Polyethylene terephthalate
		film
Haze value	35 to 45	3 to 5

Transparency results on sheets

It can be found from the results that a polyethylene terephthalate film of the present invention has 10 or more times superior transparency than a polyvinyl chloride sheet. Therefore, since polyethylene terephthalate film has good transparency and uniform surface conditions, printing sharpness, realism, etc., and it can be raised in a decorative floor covering in which patterns printed, this polyethylene terephthalate film was used in a surface layer 20.

# EXAMPLE 6

(Heat resistance test)

After installing a conventional 2.3 mm thick polyvinyl chloride resin decorative cushion floor covering, Woodleum Plus, of the COMPARATIVE EXAMPLE 1 and a 1.8 mm thick decorative floor covering of the EXAMPLE 2 on a general cement floor surface, it was measured whether damage of the product surface was shown per a period of time in the conditions that a lit cigarette and an unglazed earthenware bowl were put on each sample.

The results are represented in the following Table 2.

[Table 2] Heat resistance results per each structures

Classif	ication	A decorative A decorative		Test conditions
		floor covering of floor covering		
		COMPARATIVE EXAMPLE 2		
		EXAMPLE 1		
Lit cigarette	When left laid	Carbonization	No damages	Tested with
	parallel to a	after 30	by 5 minutes	cigarettes
	product	seconds		having a size
	When a	carbonized	No damages	of 8 mm
	cigarette is			(diameter) x 83
	crushed out			mm (length)
an unglazed ea	rthenware bowl	Damaged after	Satisfactory	Tested by
		3 seconds	until cooled	leaving boiling
				unglazed
				earthenware
				bowl on a
				product

It can be found from the results of the Table 2 that decorative floor coverings using a polyethylene terephthalate film 24 of the present invention and a polyvinyl chloride sheet substrate layer 10 containing a high content of a filler have superior heat resistance.

# **EXAMPLE 7**

(Durability(abrasion resistance) test)

An abrasion amount of a surface skin layer having a thickness of 0.25 mm in a decorative floor covering having a conventional polyvinyl chloride resin layer, that is, a Deco Tile of COMPARATIVE EXAMPLE 2 was measured and compared to an abrasion amount of a polyethylene terephthalate film layer 24 having a thickness of 0.05 mm in a decorative floor covering of EXAMPLE 2 of the present invention, using a Taber abrasion test machine in accordance with KSF 2813 (the abrasion testing method of construction materials and composing components) in order to confirm durability of a decorative floor covering of the present invention.

The results are represented in the following Table 3.

[Table 3] Abrasion resistance tests for each structure

Classification	Decorative floor covering	Decorative floor covering
	of COMPARATIVE	of EXAMPLE 2
	EXAMPLE 2	
Abrasion amout (g)	0.61	0.047
0.1 mm thickness	0.2	0.1
converted abrasion		
amount (g)		

It could be found that a decorative floor covering of the present invention was improved in durability two or more times compared to a decorative floor covering having a conventional polyvinyl chloride resin layer from the results shown in the above Table 3.

# EXAMPLE 8

(Weight comparison)

A weight per m<sup>2</sup> was 3.24 kg when a non-foaming decorative floor covering having a thickness of 2.0 mm was manufactured in COMPARATIVE EXAMPLE 3, while a weight per m<sup>2</sup> was 2.15 kg when a decorative floor covering having a thickness of 2.0 mm was manufactured in EXAMPLE 3 of the present invention, confirming that 33.6% of weight reduction was realized in a decorative floor covering of the present invention compared to a conventional non-foaming decorative floor covering.

Furthermore, insect repelling effects, heat insulating effects, a humidity controlling function, impact absorbing effects, and sound blocking effects in addition to the weight reduction can be expected from using a cork balance layer 33 in a back layer, and insect repelling effects, heat insulating effects, a humidity controlling function, and impact absorbing effects can be expected from using a wooden powder balance layer 34 in a back layer.

# **EXAMPLE 9**

15 (Sound blocking test)

After installing a 2.3 mm thick conventional vinyl cushion decorative floor covering of the COMPARATIVE EXAMPLE 1, a 1.8 mm thick conventional non-foaming vinyl cushion decorative floor covering of the COMPARATIVE EXAMPLE 3, and a 2.0 mm thick decorative floor covering of the EXAMPLE 3 on a 150 mm thick standard concrete slab, the sound blocking performance for a light floor covering impact was measured in accordance with KSF 2810 (the measuring method of floor impact sound at the building site) in order to confirm sound blocking effects of a decorative floor covering, and the measurement results are represented in the following



Table 4.

[Table 4] Sound blocking performance results on each structures

Structure	T L-	L-	Single	Improved
Structure	-	L-	Single	Improved
	class	index	evaluation	amount of
			index	sound
			(dB(A))	blocking
				performance
Standard concrete slab (150 mm)	L-75	74	75	Standard
Standard concrete slab (150 mm) + a	L-65	66.9	66.9	△8.1
decorative floor covering of				
COMPARATIVE EXAMPLE 1				
(thickness 2.3 mm)				
Standard concrete slab (150 mm) + a	L-65	66.9	66.9	△8.1
non-foaming decorative floor covering				
of COMPARATIVE EXAMPLE 3			; ;	
(thickness 1.8 mm)				
Standard concrete slab (150 mm) + a	L-50	50.1	50.1	△24.9
decorative floor covering of EXAMPLE				
3 (thickness 2.0 mm)				

In the above Table 4, L-class represents a class showing a sound blocking degree according to the Japanese Industrial Standards, L-index represents showing a sound blocking degree according to the standards prepared by Korea National Housing Corporation, and single evaluation index represents a sound blocked sound pressure level.



It could be confirmed from the above Table 4 that a decorative floor covering of the present invention had superior sound blocking effects.

# EXAMPLE 10

(Indentation test)

After applying a pressure with an indentation testing machine, an equipment of which the front end is a 6.3 mm hemispheric steel rod capable of applying a 133 N (13.6 kgf) load, for one minute in accordance with KSM 3506 (regulation on a vinyl floor sheet for a building molded of a main raw material of polyvinyl chloride resin) in order to confirm impact absorbing 10 effects of a decorative floor covering of the present invention, a indentation depth was measured by a dial gauge.

Indentation depths of a 2.3 mm thick conventional vinyl cushion decorative floor covering of the COMPARATIVE EXAMPLE 1, a 1.8 mm thick conventional polyvinyl chloride resin non-foaming decorative floor covering of 15 COMPARATIVE EXAMPLE 3, and a 2.0 mm thick decorative floor covering of EXAMPLE 3 of the present invention were measured and represented in the following Table 5.



[Table 5] Indentation results on each structures

Classification	2.3 mm thick	1.8 mm thick	2.0 mm thick
	decorative floor	decorative floor	decorative floor
	covering of	covering of	covering of
	COMPARATIVE	COMPARATIVE	EXAMPLE 3 (non-
	EXAMPLE 1	EXAMPLE 3 (non-	foaming)
	(foaming)	foaming)	
Indentation depth	1.25	0.71	0.89
(mm)			

It could be confirmed from the results of the above Table 5 that there were 25.3% improved effects compared to a 1.8 mm conventional polyvinyl 5 chloride resin non-foaming decorative floor covering although cushion characteristics were somewhat weaker than a 2.3 mm thick conventional foaming polyvinyl chloride cushion decorative floor covering.

# EXAMPLE 11

Temperature differences according to time of a 3.5 mm thick

10 conventional vinyl cushion decorative floor covering of COMPARATIVE

EXAMPLE 4 and a 2.0 mm thick decorative floor covering of EXAMPLE 3 of
the present invention were measured and represented in the following Table

6.



[Table 6] Heat accumulating effects on each structures

Time lapsed	3.5 mm thick	vinyl cushion	2.0 mm thick d	ecorative floor
	flooring of CC	MPARATIVE	covering of E	EXAMPLE 3
	EXAMPLE 4			
	Temperature	Temperature	Temperature	Temperature
	(℃)	difference	(℃)	difference
		(℃)		(℃)
Product initial	54	-	49	<del>-</del>
temperature	42	12	39	10
l hour	35	19	31	18
2 hours	31	23	27	22
3 hours	27	27	25	24
4 hours				

It could be confirmed from heat accumulating effect results on each structure of the above Table 6 that a 2.0 mm thick decorative floor covering of the present invention had 3 °C or more heat insulating effects than a 3.5 mm thick conventional vinyl cushion decorative floor covering.

A decorative floor covering of the present invention is a decorative floor covering which has superior surface heat resistance, printing sharpness and realism, and it can be installed conveniently since not only can non-floamed polyvinyl chloride resin be used as a substrate layer, but also the weight of the floor covering is light when a light weight back layer is comprised under the substrate layer.

While the present invention has been described in detail with



# PCT/KR00/00664

reference to the preferred embodiments, those skilled in the art will appreciate that various modifications and substitutions can be made thereto without departing from the spirit and scope of the present invention as set forth in the appended claims.



## WHAT IS CLAIMED IS:

- 1. A decorative floor covering downwardly comprising a surface layer, and a polyvinyl chloride resin substrate layer, wherein the surface layer comprises a polyethylene terephthalate film layer.
- 2. A decorative floor covering in accordance with claim 1, wherein the polyethylene terephthalate film layer is a film in which a primer selected from the group consisting of polyvinyl acetate based, polyurethane acrylate based, and ethylene vinyl acetate based primers is coated to a film thickness of from 0.1 to 10  $\mu$ m on top and bottom sides of the polyethylene terephthalate film layer.
  - 3. A decorative floor covering in accordance with claim 1, wherein a thickness of the polyethylene terephthalate film layer is from 10 to 100  $\mu$ m.
- 4. A decorative floor covering in accordance with claim 1, wherein the polyvinyl chloride resin substrate layer is a non-foamed polyvinyl chloride sheet comprising 100 weight parts of polyvinyl chloride resin, 50 to 400 weight parts of one or more inorganic fillers selected from the group consisting of calcium carbonate, talc, wollastonite, and silica, or 5 to 30 weight parts of one or more metallic powders selected from the group consisting of aluminum, copper, and iron as fillers.
  - 5. A decorative floor covering in accordance with claim 4, wherein the polyvinyl chloride resin substrate layer 10 is a non-foamed polyvinyl chloride sheet manufactured by kneading a polyvinyl chloride resin composition comprising 100 weight parts of polyvinyl chloride resin, a plasticizer of 30 to

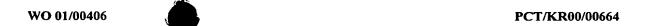


50 weight parts of dioctyl phthalate, heat resistant stabilizers of 3 to 5 weight parts of barium-zinc based compound and 3 to 5 weight parts of epoxy compound, and fillers of 50 to 400 weight parts of calcium carbonate (CaCO<sub>3</sub>) and 3 to 5 weight parts of pigment in a rolling mill having a 5 temperature of 160 to 190 °C, and pressure rolling it to a thickness of 0.8 to 1.3 mm.

- 6. A decorative floor covering in accordance with claim 1, wherein the polyvinyl chloride resin substrate layer is comprised of glass fiber scrim, or woven or nonwoven glass fiber.
- 7. A decorative floor covering in accordance with claim 1 further comprising a light back layer beneath the substrate layer.
- 8. A decorative floor covering in accordance with claim 7, wherein the light back layer comprises one or more layers selected from the group consisting of a cork balance layer, a wooden powder balance layer, and a fiber layer.
- 9. A decorative floor covering in accordance with claim 8, wherein the cork balance layer is a sheet which is cut to a thickness of 1.0 to 2.0 mm with a slicing machine after a cork layer of a natural tree is peeled off, pulverized to a particle size of 5 to 10 meshes, put into a large cylinder together with adhesive, and bonded by applying pressure.
  - 10. A decorative floor covering in accordance with claim 8, wherein the wooden powder balance layer is a sheet which is rolled to a thickness of 0.8 to 1.0 mm using a two sets of pressure rolls after adding 100 to 150 weight parts of wooden powder having a particle size of 200 to 300 meshes,

3 to 5 weight parts of heat resistant stabilizer barium-zinc based compound,
1 to 3 weight parts of internal antiadditive stearic acid, 30 to 50 weight parts
of plasticizer dioctyl phthalate, and 3 to 5 weight parts of pigment to 100
weight parts of polyvinyl chloride resin, and kneading all the components
together.

- 11. A decorative floor covering in accordance with claim 8, wherein the fiber layer is selected from the group consisting of woven or unwoven polyester, woven or unwoven polypropylene, and woven or unwoven glass fiber, the woven thickness being from 10s x 10s to 25s x 15s, and the density being from 20 x 20 to 30 x 30 roll/inch.
  - 12. A decorative floor covering in accordance with claim 1 comprising:
    - i ) a surface treated layer;
    - ii) a polyethylene terephthalate film layer:
- 15 iii) a polyvinyl chloride resin intermediate layer:
  - iv) a polyvinyl chloride resin substrate laver:
  - v) an adhesive layer; and
  - vi) a light back layer.
- 13. A decorative floor covering in accordance with claim 12, wherein 20 the surface treated layer of i) is a layer in which a surface treated layer composition comprising a resin selected from the group consisting of urethane acrylate, silicone acrylate, and epoxy acrylate is coated on the polyethylene terephthalate film layer, and cured with ultraviolet rays.



14. A decorative floor covering in accordance with claim 13, wherein the surface treated layer composition further comprises an acryl based or urethane based bead having a particle size of 5 to 20  $\mu$ m.

- 15. A decorative floor covering in accordance with claim 12, wherein the polyvinyl chloride resin intermediate layer of ii) is a sheet having a thickness of 0.1 to 1 mm manufactured by pressure rolling in a calender a polyvinyl chloride resin composition comprising 100 parts of polyvinyl chloride resin, 25 to 50 weight parts of dioctyl phthalate, 50 to 150 weight parts of calcium carbonate, 3 to 5 weight parts of titanium oxide, and 2 to 5 weight parts of heat stabilizer.
  - 16. A decorative floor covering in accordance with claim 12, wherein the total thickness of the i) surface treated layer, ii) polyethylene terephthalate film layer, iii) polyvinyl chloride resin intermediate layer, iv) polyvinyl chloride substrate layer, and v) adhesive layer is from 1 to 1.5 mm.
  - 17. A decorative floor covering in accordance with claim 12, wherein the adhesive layer of v) is a sheet in which urethane resin, acrylic resin or vinylacetate resin is rolled to a thickness of 0.03 to 0.2 mm using 4 calender rolls at a temperature of 80 to 160  $^{\circ}$ C, an impregnated sheet, or a plastisol coating gel.

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20 18. A method for manufacturing a decorative floor covering comprising a surface layer downwardly comprising a surface treated layer, a polyethylene terephthalate film layer, and a polyvinyl chloride resin intermediate layer, and a polyvinyl chloride resin substrate layer comprising

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the steps of:

- a) manufacturing a polyvinyl chloride resin substrate layer;
- b) manufacturing a pressed sheet by laying up a polyvinyl chloride resin intermediate layer on the substrate layer of step a), applying pressure, and pressing;
- c) manufacturing a half-finished sheet by laying up a polyethylene terephthalate film layer under which a certain pattern is printed on the pressed sheet of step b), applying pressure, and pressing; and
- d) forming a surface treated layer by coating a surface treated layer composition on the half-finished sheet of step c) and UV (ultraviolet ray) curing.
- 19. A method for manufacturing a decorative floor covering in accordance with claim 18, wherein the pressing of step b) is carried out after preheating a substrate layer to a temperature of 140 to 170 ℃ before laying up a polyvinyl chloride resin intermediate layer.
  - 20. A method for manufacturing a decorative floor covering in accordance with claim 18, wherein the pressing of step c) is carried out after preheating a pressed sheet to a temperature of 140 to 170 ℃ before laying up a polyethylene terephthalate film layer 24.
- 21. A method for manufacturing a decorative floor covering in accordance with claim 18, wherein the polyethylene terephthalate film layer of step c) is a film in which a primer selected from the group consisting of polyvinyl acetate based, polyurethane acrylate based, and ethylenevinyl acetate based primers is coated to a film thickness of 0.1 to 10 [All] on top



and bottom sides of the polyethylene terephthalate film layer.

- 22. A method for manufacturing a decorative floor covering in accordance with claim 18, wherein the polyethylene terephthalate film layer of step c) has a thickness of 10 to 100  $\mu$ m.
- 23. A method for manufacturing a decorative floor covering in accordance with claim 18, wherein the polyvinyl chloride resin substrate layer of step a) is manufactured by kneading a polyvinyl chloride resin composition comprising 100 weight parts of polyvinyl chloride resin, 50 to 400 weight parts of one or more inorganic fillers selected from the group consisting of 10 calcium carbonate, talc, wollastonite, and silica, or 5 to 30 weight parts of one or more metallic powders selected from the group consisting of aluminum, copper, and ion as fillers in a rolling mill of 160 to 190 ℃, and rolling to a thickness of 0.8 to 1.3 mm.
- 24. A method for manufacturing a decorative floor covering 15 comprising a surface layer downwardly comprising a surface treated layer, a polyethylene terephthalate film layer, and a polyvinyl chloride resin intermediate layer, a polyvinyl chloride resin substrate layer, and a light back layer comprising the steps of:
  - a) manufacturing the first polyvinyl chloride resin substrate layer;
- 20 b) pressing while applying pressure in a embossing roll after positioning a glass fiber scrim, or a woven or unwoven glass fiber underneath the first substrate layer 10 under the condition that a surface of the first polyvinyl chloride resin substrate layer 10 of step a) maintains latent heat of 100 to 150 ℃:

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- c) manufacturing the second polyvinyl chloride resin substrate layer;
- d) coating a plastisol underneath the second polyvinyl chloride resin substrate layer, positioning a light back layer of a fiber layer, applying pressure, and pressing with a press roll at a heating drum at 140 to 150 ℃;
- e) manufacturing a polyvinyl chloride resin intermediate layer;
- f) manufacturing a pressed sheet by applying pressure at a press after downwardly positioning a polyvinyl chloride resin intermediate layer, the first polyvinyl chloride resin substrate layer under which glass fiber scrim, or woven or unwoven glass fiber is pressed, the second polyvinyl chloride resin substrate layer under which a light back layer of a fiber layer is pressed, and preheating to a temperature of 140 to 170 °C;
- g) manufacturing a half-finished sheet by applying pressure and pressing at an embossing roll after preheating the pressed sheet of step f) to a temperature of 140 to 170 °C and positioning a polyethylene terephthalate film layer on a polyvinyl chloride resin intermediate layer; and
- h) forming a surface treated layer by UV (ultraviolet rays) curing after coating a surface treated layer composition on the half-finished sheet of step g).
- 25. A method for manufacturing a decorative floor covering in accordance with claim 24, wherein the polyethylene terephthalate film layer of step g) is a film in which a primer selected from the group consisting of



polyvinyl acetate based, polyurethane acrylate based, and ethylenevinyl acetate based primers is coated to a film thickness of 0.1 to 10  $\mu$ m on top and bottom sides of the polyethylene terephthalate film layer.

- 26. A method for manufacturing a decorative floor covering in accordance with claim 24, wherein the polyethylene terephthalate film layer of step q) has a thickness of 10 to 100  $\mu$ m.
- 27. A method for manufacturing a decorative floor covering in accordance with claim 24, wherein the first polyvinyl chloride resin substrate layer 10 of step a) and the second polyvinyl chloride resin substrate layer of step c) are manufactured by kneading a polyvinyl chloride resin composition comprising 100 weight parts of polyvinyl chloride resin, 50 to 400 weight parts of one or more inorganic fillers selected from the group consisting of calcium carbonate, talc, wollastonite, and silica, or 5 to 30 weight parts of one or more metallic powders selected from the group consisting of aluminum, copper, and iron as fillers in a rolling mill at 160 to 190 °C, and rolling to a thickness of 0.8 to 1.3 mm.
- 28. A method for manufacturing a decorative floor covering comprising a surface layer downwardly comprising a surface treated layer, a polyethylene terephthalate film layer, and a polyvinyl chloride resin intermediate layer, a polyvinyl chloride resin substrate layer, and a light back layer comprising the steps of:
  - a) manufacturing a polyvinyl chloride resin substrate layer;
  - b) laying up a polyvinyl chloride resin intermediate layer on the preheated substrate layer of step a), applying pressure, and

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pressing;

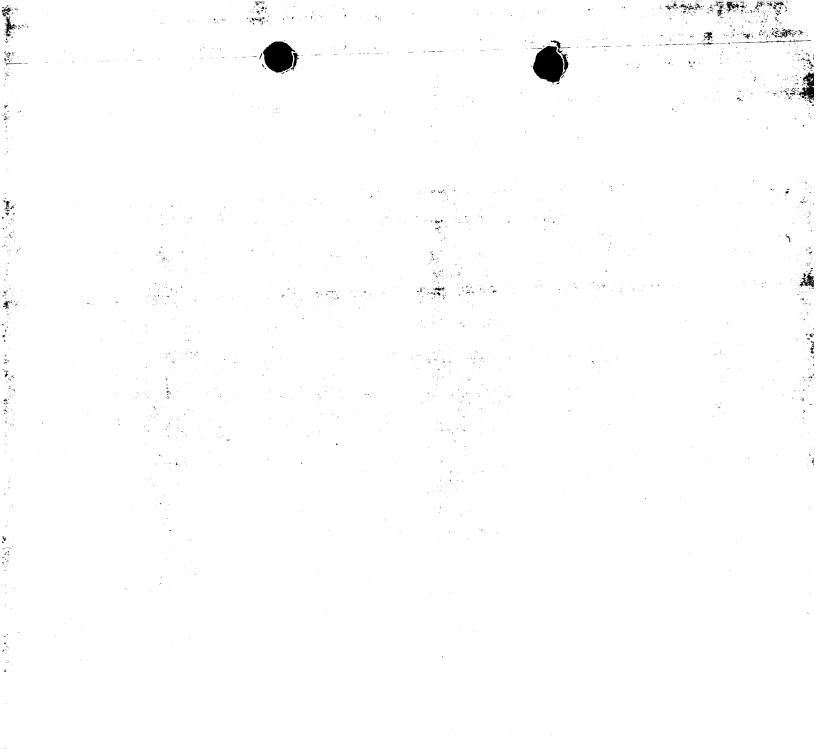
- c) manufacturing a half-finished sheet by laying up a polyethylene terephthalate film layer on the preheated polyvinyl chloride resin intermediate layer of step b), applying pressure, and pressing:
- d) laying up an adhesive layer underneath the preheated halffinished sheet of step c), applying pressure, and pressing;
  - e) laying up a light back layer underneath the adhesive layer of step d), applying pressure, and pressing; and
- f) forming a surface treated layer after coating a surface treated layer composition on the pressed polyethylene terephthalate film layer of step e), and curing.
- 29. A method for manufacturing a decorative floor covering in accordance with claim 28, wherein the laying up and pressing of the light back layer is laying up and pressing a fabric layer underneath a wooden powder balance layer after laying up and pressing a cork balance layer underneath an adhesive layer, or laying up and pressing a wooden powder balance layer underneath an adhesive layer.
- 30. A method for manufacturing a decorative floor covering in accordance with claim 28, wherein the polyethylene terephthalate film layer of step c) is a film in which a primer selected from the group consisting of polyvinyl acetate based, polyurethane acrylate based, and ethylenevinyl acetate based primers is coated to a film thickness of 0.1 to 10 \(mu\) on top and bottom sides of the polyethylene terephthalate film layer.
  - 31. A method for manufacturing a decorative floor covering in

accordance with claim 28, wherein the polyethylene terephthalate film layer of step c) has a thickness of 10 to 100  $\,\mu\mathrm{m}$ .

32. A method for manufacturing a decorative floor covering in accordance with claim 28, wherein the polyvinyl chloride resin substrate layer of step a) is manufactured by kneading a polyvinyl chloride resin composition comprising 100 weight parts of polyvinyl chloride resin, 50 to 400 weight parts of one or more inorganic fillers selected from the group consisting of calcium carbonate, talc, wollastonite, and silica, or 5 to 30 weight parts of one or more metallic powders selected from the group consisting of aluminum, 10 copper, and iron as fillers in a rolling mill of 160 to 190 ℃, and rolling to a thickness of 0.8 to 1.3 mm.

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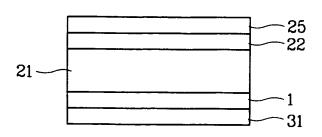
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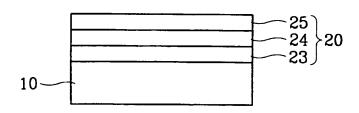


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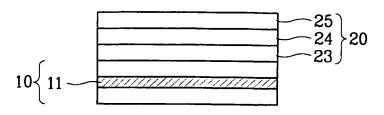
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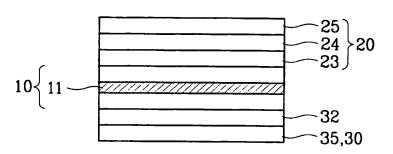


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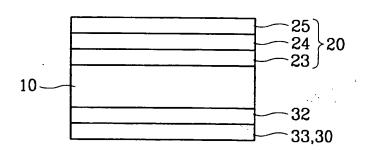


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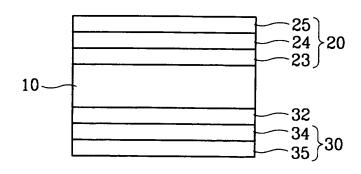
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International application No. PCT/KR00/00664

### CLASSIFICATION OF SUBJECT MATTER

IPC7 B32B 27/18

According to International Patent Classification (IPC) or to both national classification and IPC

Minimun documentation searched (classification system followed by classification symbols)

IPC7 B32B 27/08, B32B 27/00, B32B 33/00

Documentation searched other than minimum documentation to the extent that such documents are included in the fileds searched Korean Patents and Applications for Invention since 1975

Korean Utility Models and Applications for Utility Models since 1975

Electronic data base consulted during the intertnational search (name of data base and, where practicable, search trerms used) NPS, PAJ

#### DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
А	JP 08-174783 A (Dainippon Printing Co., Ltd.) See Claim1-9, Fig.1-2	1
A	KR 97-33802 A (Hyundai Motor Co., Ltd.) See Abatract, Claim1-3	1
A	KR 94-13816 A (Lucky Co., Ltd.) See Abatract. Claim1-10, Fig.1-4	1

Further documents are listed in the continuation of Box C.	See patent family annex.
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- Special categories of cited documents:
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Date of the actual completion of the international search Date of mailing of the international search report 09 OCTOBER 2000 (09.10.2000) 10 OCTOBER 2000 (10.10.2000)

Name and mailing address of the ISA/KR Korean Industrial Property Office

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# INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No. PCT/KR00/00664

Publication date	Patent family member(s)	Publication date	Patent document cited in search report	
	None	09.07.96	JP 08-174783	
	None	22.07.97	KR 97-33802	:
	None	16.07.94	KR 94-13816	
	None	22.07.97	KR 97-33802	